Final Remedial Investigation/Feasibility Study Work Plan

Bremerton Gas Works Superfund Site

Prepared for: Cascade Natural Gas Corporation

Aspect Project No. 080239-005 • Anchor QEA Project No. 131014-01.01 February 28, 2017

Prepared by



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- H Site Health and Safety Plan (Anchor QEA, LLC)

Abbreviations

Anchor QEA, LLC

AOC Administrative Settlement Agreement and Order on Consent for Remedial

Investigation Feasibility Study

ARAR applicable or relevant and appropriate requirement

Aspect Aspect Consulting, LLC
AST aboveground storage tank
bgs below ground surface

BTAG Biological Technical Assistance Group BTEX benzene, toluene, ethylbenzene, and xylenes

Cascade Natural Gas Corporation

CERCLA Comprehensive Environmental Response, Compensation, and Liability Act

CFR Code of Federal Regulations

City City of Bremerton contaminant of concern

COPC contaminant of potential concern

cPAH carcinogenic polycyclic aromatic hydrocarbon

CSM conceptual site model CSO combined sewer overflow

CWA Clean Water Act

DNAPL dense non-aqueous phase liquid

DNR Washington State Department of Natural Resources

DQO data quality objective

DU data usability

E&E Ecology & Environment, Inc.

Ecology Washington State Department of Ecology

EcoSSL ecological soil screening level EDD electronic data deliverable

EM electromagnetic

ENVVEST Environmental Investment Project
EPA U.S. Environmental Protection Agency

ERA ecological risk assessment

ER-L effect range-low ER-M effect range-medium FS Feasibility Study

GPR ground-penetrating radar HHRA human health risk assessment

HPAH high-molecular-weight polycyclic aromatic hydrocarbon

ISA initial study area

KPHD Kitsap Public Health District

Lent's Lents and Blombergs

LNAPL light non-aqueous phase liquid

LPAH low-molecular-weight polycyclic aromatic hydrocarbon

MCL maximum contaminant level MDAC minimum data acceptability criteria

mg/kg milligram(s) per kilogram
MGP manufactured gas plant

MTCA Washington State Model Toxics Control Act

NAPL non-aqueous phase liquid

NOAA National Oceanographic and Atmospheric Administration

NRHP National Register of Historic Places

Order Administrative Order for a Pollution Incident

PAH polycyclic aromatic hydrocarbon

PCB polychlorinated biphenyl PCP pentachlorophenol

PHS Priority Habitats and Species Program

PQL practical quantitation limit PRG preliminary remediation goal

PSAMP Puget Sound Assessment and Monitoring Program

RAO remedial action objectives

RCRA Resource Conservation and Recovery Act

RI Remedial Investigation RSL regional screening level SCO sediment cleanup objective

Site Bremerton Gas Works Superfund Site SMS Sediment Management Standards

SOW Statement of Work

SPME solid-phase microextraction

SQAPP Sampling and Quality Assurance Project Plan

SVOC semivolatile organic compound TBA Targeted Brownfields Assessment

TBC to be considered

TCRA time critical removal action

TEQ toxic equivalent

TMDL total maximum daily load TOC total organic carbon

TPH total petroleum hydrocarbons

Tribe Suquamish Tribe

TS total solids

 $\begin{array}{ll} \mu g/kg & \text{microgram(s) per kilogram} \\ \mu g/L & \text{microgram(s) per liter} \\ USCG & U.S. \ Coast \ Guard \end{array}$

UST underground storage tank

UV ultraviolet

VOC volatile organic compound

WDFW Washington Department of Fish and Wildlife

WDOH Washington Department of Health

1 Introduction

Cascade Natural Gas Corporation (Cascade) is conducting a Remedial Investigation (RI) and Feasibility Study (FS) at the Bremerton Gas Works Superfund Site (Site) under the direction of the U.S. Environmental Protection Agency (EPA). This Final Remedial Investigation/Feasibility Study Work Plan (Work Plan) presents detailed descriptions of the procedures and activities to be performed to complete the RI/FS. This Work Plan was prepared as required by the Administrative Settlement Agreement and Order on Consent for Remedial Investigation Feasibility Study (AOC; EPA, 2013a) and accompanying Statement of Work (SOW) for the Bremerton Gas Works Superfund Site.

The Site, whose boundaries have not yet been defined, encompasses approximately 2.8 acres of industrial upland property and marine beachfront on the south shore of the Port Washington Narrows in Bremerton, Kitsap County, Washington. The Site location is depicted on Figure 1-1.

As an initial step in the RI/FS process, Cascade prepared a Final Scoping Memorandum (Aspect and Anchor, 2015; Scoping Memorandum) to identify the tasks necessary to conduct and complete the RI/FS. This Work Plan documents decisions and evaluations made during the scoping process and presents anticipated future tasks to complete the RI/FS. Following finalization of this Work Plan, a series of marine and upland field investigations will be conducted to gather data relevant to the Site. ¹

The data collected during field investigations will be used to develop the Conceptual Site Model (CSM) for the Site, which will be presented together with the data in an RI Report. Concurrent with the RI Report, Cascade will prepare baseline risk assessments. Following finalization of the RI Report and the baseline risk assessments, a FS Report will be prepared that develops and evaluates potential remedies for the Site. EPA will use this information to develop a Proposed Plan. The phases of the RI/FS tasks are shown in the RI/FS Work Flow Chart (Figure 1-2).

A manufactured gas plant (MGP) formerly operated on a portion of the Site. Other historical uses on or near the Site include bulk petroleum storage and distribution, equipment storage, boat maintenance, metal fabrication, and automobile salvage. Previous investigations have identified elevated concentrations of hazardous substances in soil, groundwater, and sediments, attributable to these historical activities. Currently, portions of the Site are largely vacant and unused.

In accordance with the AOC and SOW, this Work Plan includes detailed sampling and quality assurance project plans. The Sampling and Quality Assurance Project Plan (SQAPP) for the upland and marine portions of the Site are included as Appendices A and B, respectively.

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¹This Work Plan has been produced based on the current CSM which was developed using available information. The Cascade and EPA project teams may agree that elements of this Work Plan should change based on data collected during the RI. EPA must approve all changes to this Work Plan.

1.1 Objectives of the RI/FS

The objectives the RI/FS for the Site are the following:

- 1. Investigate and define physical, chemical and biological characteristics of the Site;
- 2. Define the sources, nature, and distribution of contaminants;
- 3. Provide sufficient information to calculate and assess the current and future potential risks to human health and the environment; and
- 4. Provide sufficient information to identify and evaluate remedial alternatives, conceptually design the remedial alternatives, and select a remedy.

The RI/FS will be conducted in accordance with the provisions of the AOC, SOW, the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA), the National Contingency Plan, and EPA guidance, including, but not limited to, Guidance for Conducting Remedial Investigations and Feasibility Studies Under CERCLA (EPA 1988a), and Guidance for Data Usability in Risk Assessment (EPA 1992).

Work undertaken throughout the RI/FS process will be conducted in coordination with key stakeholders and the public, with EPA serving in a lead role in those coordination efforts (see Section 9). The work will be conducted, not only in accordance with the legal requirements mentioned previously, but also consistent with other applicable EPA policies and practices, such as EPA's responsibility to consult with the Suquamish Tribe (Tribe) on a government-to-government basis, EPA's Tribal Policy, and EPA's guiding principles on Environmental Justice.

1.2 Work Plan Organization

This Work Plan is organized into the following Sections:

- Section 2 Background and Setting. Section 2 provides a description of the Site location; a summary of known current and historical uses of the Site and adjacent properties and aquatic lands; a summary of the Site environmental setting including regional and Site geology and hydrogeology; a discussion of current demographics and land use; a summary of the characteristics of the Port Washington Narrows; and a description of natural and cultural resources in the Site vicinity.
- Section 3 Initial Evaluation. Section 3 presents the regulatory requirements and provides a summary of the previous work conducted that is relevant to the RI/FS, including previous Site investigations, previous removal actions, and available existing data. A summary of the existing data for soil, groundwater, and sediment is also presented in this section.
- Section 4 Preliminary Conceptual Site Model. Section 4 presents a conceptual understanding of the Site based on the information discussed in Sections 2 and 3, including a summary of the contaminants of potential concern, their sources, transport mechanisms, exposure pathways and receptors.
- Section 5 RI/FS Approach. Section 5 presents the approach for completing the RI/FS and the rationale behind the approach, including identification of the data needs, a

- summary of the risk assessment approach, a general discussion of the components of the upland and marine portions of the RI/FS, and potential contingent actions.
- Section 6 RI Tasks. Section 6 presents a summary of the tasks to be conducted for completion of the RI.
- Section 7 FS Planning. Section 7 presents a summary of the tasks to be conducted for completion of the FS. It also includes a discussion of potentially applicable remedial technologies for the Site, a summary of remedial approaches that have been implemented at similar sites, and the data needed to develop and evaluate remedial alternatives for the Site.
- Section 8 Schedule. Section 8 presents the schedule for completion of the RI/FS including a field data collection schedule and the general schedule for subsequent tasks and reports.
- Section 9 Project Management Plan. Section 9 presents the project management plan, including a data management plan.
- Section 10 References. Section 10 presents a list of the references cited within this Work Plan.

2 Background and Setting

This Section describes the property upon which the former gas works was located and the properties surrounding the former gas works and discusses the operational and regulatory history of those properties.

2.1 Site Location and Description

The former gas works was located between Thompson Drive and Pennsylvania Avenue (Figure 2-1) on approximately 2.8 acres of property along the south shore of Port Washington Narrows in Bremerton, Washington. The historical street addresses for the former gas works included 1720 and 1800 Thompson Drive.

The real property upon which the former gas works was located (Former Gas Works Property) relative to current parcel boundaries is shown on Figure 2-1. Due to a boundary line adjustment in 1992, the Former Gas Works Property includes portions of two existing tax parcels:

- **Kitsap County Parcel No. 3711-000-0010-0409 (McConkey Property).** This parcel is owned by the McConkey Family Trust. The former gas works covered the entire parcel. No current or historical street address has been identified for this parcel.
- Kitsap County Parcel No. 3741-000-022-0101 at 1701 Pennsylvania Avenue (Sesko Property). This parcel is owned by Natasha Sesko. The former gas works covered the northwestern portion of this parcel.

The following properties are located near the Former Gas Works Property and have had either suspected or confirmed releases of contaminants from historical operations unrelated to the former gas works:

- 1723 Pennsylvania Avenue (Penn Plaza Property). This property is owned by Penn Plaza Storage, LLC. There are multiple street addresses associated with this property, but it is listed in the Kitsap County assessor's database as 1723 Pennsylvania Avenue.
- 1701 Thompson Drive (Former ARCO Property). This property is owned by Pipeworks Mechanical & Service, Inc. It is located southwest of the Former Gas Works Property, across Thompson Drive.
- 1702 Pennsylvania Avenue (Former SC Fuels Property). This property is owned by NFS Properties 2, LLC. It is located east of the Sesko Property, across Pennsylvania Avenue.

The Port Washington Narrows is located north of the McConkey, Sesko, and Former SC Fuels Properties. The Port Washington Narrows consists of aquatic lands owned by the State of Washington and managed by the Washington State Department of Natural Resources (DNR).

2.2 Site Uses Prior to 1930

The Port Washington Narrows and the adjacent uplands are located in the traditional territory of the Suquamish Tribe, a Southern Coast Salish community speaking a dialect of the Southern Lushootseed language (Suttles and Lane, 1990). Shoreline locations in Dyes Inlet would have been available after stabilization of sea levels in the mid-Holocene (Thorson, 1980); therefore, Native American use of the area may date back 10,000 years. A variety of traditional activities took place in the general vicinity. In 1855, the Tribe signed the Treaty of Point Elliott, which ceded lands and established the reservation at Port Madison. The Tribe retained "the right of taking fish at usual and accustomed grounds and stations" (Treaty of Point Elliott, 1855), and the Port Washington Narrows is within the Tribe's adjudicated Usual and Accustomed area.

2.3 Current and Historical Use and Operations

Historical use and operations on the properties and aquatic lands are based on historical records, including aerial photographs, interviews with current and former workers, owners, area residents, historical maps, deeds, Washington State Department of Ecology (Ecology) records, City of Bremerton (City) records, and DNR lease records. A number of historical documents are included in previous assessments of historical Site use (TechLaw, 2006; Hart Crowser, 2007). Available and relevant historical records are provided in Appendix C for reference.

Historical and current operations on the Former Gas Works Property (which consists of the entire McConkey Property and a portion of the Sesko Property) as well as historical and current operations on the other portion of the Sesko Property are described in Section 2.3.1. Historical and current operations on adjoining properties are described in Section 2.3.2.

2.3.1 Operations on McConkey and Sesko Properties

2.3.1.1 Former Gas Works Operations

In 1930, the Former Gas Works Property was developed as a gas works (a.k.a., manufactured gas plant, or MGP). Gas works were a common industry in large and small towns throughout the United States and Europe from approximately the mid-1800s to the mid-1900s. At a gas works, coal, coke, and/or petroleum products were heated in furnaces to produce manufactured gas, which was subsequently distributed via a gas piping network to the surrounding homes and businesses for heating, cooking, and lighting. Gas works used or generated several products and byproducts, including non-aqueous phase liquids (NAPLs) such as oils and tars, aqueous waste streams, and solid materials containing chemicals that may pose a risk to human health or the environment because they are toxic or carcinogenic (resulting in cancer effects). These contaminants include hydrocarbons such as benzene, toluene, ethylbenzene, and xylenes (BTEX) and polycyclic aromatic hydrocarbons (PAHs), which can persist for a long time in the environment. Contaminant releases from historical gas works operations at other locations have resulted in sites where contamination remains in the subsurface as NAPLs, sorbed to soil or sediments or dissolved in the groundwater.

Because of the potential hazards posed by historical gas works facilities, these facilities are often the focus of state-led or federally led efforts to investigate and clean up contamination to protect human health and the environment. To characterize and remediate these facilities, it is important to understand traditional gas works operations, the types of contaminants that may be present, and where contaminants may have been released. This Section provides a summary of what is known about operations at the former gas works based on historical documentation and what is assumed based on typical gas works operations. This Section also identifies the contaminants usually associated with gas works feedstocks, fuels, and byproducts that may be present at the Site. Uncertainties about historical practices and potential releases will be addressed through field investigations as described in this Work Plan. Further discussion of potential release mechanisms and transport of contaminants in the subsurface is provided in Section 4, Preliminary Conceptual Site Model.

The operational history of the former gas works is as follows:

- **1930 to 1931.** The former gas works was constructed by the Western Gas and Utilities Corporation (Western).² It included a dock on aquatic lands initially leased from the DNR on November 25, 1930 (Former Gas Works Dock).
- 1931 to 1955. Manufactured gas was produced using the carbureted water-gas process, from feedstocks of coal, coke briquettes, and petroleum products. In the 1940s, a standby plant for producing natural gas by blending liquefied petroleum (butane or propane) and air was installed. Gas produced at the Former Gas Works Property in the 1940s and 1950s was from manufactured gas and from butane-air. In 1952, the Former Gas Works Property was transferred from Western to Bremerton Gas Company, and in 1953 it was transferred to Cascade. In approximately 1955 (Simonson, 1997b), manufactured gas operations ceased, and all gas was produced from butane-air mixing.
- 1955 to 1963. Natural gas was produced from butane-air mixing. In 1963, with the completion of a natural gas pipeline to the region, gas production ceased.
- 1963 to 1972. Some of the structures and tanks were removed between 1964 and 1965, and the concrete piers supporting the tanks were jackhammered and hauled away (White 1998). The former plant building was reportedly used for pipe storage and, for a short time, magnesium mining research (*Bremerton Sun*, 1972). In 1972, the remaining structures, including the former plant building, were sold and dismantled.

In 1972, the Former Gas Works Property was acquired by Harold D. and L. Irene Lent and Theodore and Marian J. Blomberg, doing business as "Lent, Blomberg, Lent." The Lent and Blomberg families operated several businesses near the Former Gas Works Property,

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²In 1931, the Western Gas and Utilities Corporation changed its name to the Western Gas Company of Washington. The Western Gas and Utilities Corporation and the Western Gas Company of Washington are collectively referred to herein as "Western."

³Typically, diesel-range fuel oils were used for petroleum feedstock for the carbureted water-gas process (Hatheway, 2012). However, one historical map (Sanborn, 1946) indicates that gasoline and fuel oil were stored in the northeast corner of the Former Gas Works Property.

including an oil distribution business on the Sesko Property under the name Lents, Inc. (see further discussion in Section 2.3.1.3). All entities and individuals associated with the Lents and Blombergs are referred to in this Work Plan as "Lent's."

In 1979, Paul and Margaret McConkey acquired most of the Former Gas Works Property. The McConkeys acquired the remainder of the Former Gas Works Property in 1985. A portion of the Former Gas Works Property was sold to William Sesko in 1992.

The summary of gas works operations provided in this Section combines available historical information about the layout and operations of the former gas works with information compiled from multiple sources regarding the operations of typical manufactured gas facilities, including generated byproducts and likely sources of releases of hazardous substances. Whereas this summary provides an overview of operations at the former gas works, it likely does not provide a complete picture of all sources, disposal areas, and spills and/or releases that may have occurred, which will be investigated primarily through the collection and evaluation of data as described in this Work Plan. Chemical feedstocks and potential byproducts typical of carbureted water-gas production include the following:

- Feedstock and Fuels: Gasoline and Diesel Fuel Oil, Coal, or Coke Briquettes. The contaminants potentially associated with feedstock and fuels include the following:
 - o BTEX;
 - o Naphthalenes; and
 - o PAHs.
- Byproducts: Light Oil, Carbureted Water-Gas Tar, Ash, Clinker, Slag, Soot, and Spent Purifier Filter Media. The contaminants potentially associated with byproducts include the following:
 - o BTEX;
 - Naphthalenes;
 - o PAHs;
 - Phenols;
 - Other semivolatile organic compounds (SVOCs), including creosol, carbazole, and dibenzofuran.

Section 4.4 provides further discussion of the Site-specific COPCs.

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⁴ Two byproducts typically generated at coal and/or oil gas plants, ammoniacal liquor and lampblack (carbon soot), were generally not generated in significant quantities by the carbureted water-gas process (Hatheway, 2012).

Production of natural gas using liquefied petroleum (butane or propane) blended with air is not anticipated to have resulted in contamination of the subsurface because butane and propane are gases at atmospheric conditions.

A flow chart showing the gas works process as understood at the Site (based on available plant maps and typical carbureted water-gas operations), including the production of byproducts, is presented on Figure 2-2. The locations of key plant features are shown on Figure 2-3. The general sequence of operations is as follows:

- Product Delivery and Storage. Solid feedstocks (coal and coke briquettes) were transported to the Site by barge and offloaded via a winch to a storage slab located in the northwest corner of the Former Gas Works Property. Petroleum products were also delivered to the former gas works via barge and conveyed via a pipeline up the Former Gas Works Dock to storage tanks located in the northeast corner of the Former Gas Works Property.
- Gas Generation and Purification. These operations were located in the north-central portion of the Former Gas Works Property (Figure 2-3). Two generator sets (furnaces) were located in the main plant building: one in the northern portion of the building and one in the middle of the building (Simonson, 1997b). The main plant building had a concrete floor (Simonson, 1997b). Coal and coke were placed in the generators and heated, and fuel oil was sprayed into the generators to produce gas. The resulting gas stream was then passed through a series of devices to cool the gas and remove impurities. These devices are described below:
- **Scrubber.** After gas generation comes clarification, in which tar is separated from the gas using a scrubber or similar equipment. These devices are typically located adjacent to the generator sets. A historical plant map shows the scrubber located directly west of the generator sets. A former plant worker indicated that the scrubber consisted of a tank with wooden slots and water to "wash out" the gas (Simonson, 1997b). An engineer's report (Tymstra, 1942) indicates that wood chips and excelsior (i.e., wood shavings) were used to remove tar from the gas.

The clarification process typically produced tar, tar-soaked wood chips or shavings, gas liquor (aqueous solutions containing dissolved and suspended tar particles), and tar-water emulsions. Light oils may also have been produced in the scrubbing process. Tar-water emulsions from scrubbers were typically removed from clarification equipment and transported to residual management areas to separate tar from the water (Hatheway, 2012). The fate of byproducts and residuals is discussed in the bullet "Residuals Management."

- O Gas Holder. A large gas holder was located south of the scrubber, west of the main plant building. The bottom of the gas holder was reportedly 15 feet deep and contained tar and water (Simonson, 1997a). The materials used to construct the base of the gas holder are unknown.
- O Purifier. Gas was passed through a bed of filter media to remove impurities such as sulfide from the gas. Typical filter media included wood chips and/or iron oxide. An engineer's report (Tymstra, 1942) indicated that iron-oxide-covered chips were used at the gas works to remove sulfur compounds from gas. Multiple purifiers in parallel were typically installed to allow changeout of purifier media without interrupting the process

(Hatheway, 2012). Three purifiers were located at the Former Gas Works Property south of the large gas holder. In addition to the generation of spent purifier media, which included some accumulated tar (Tymstra, 1942), some liquid streams (including tar, gas liquor, and light oil) may have condensed during purification and were typically manually removed from the purifier box (Hatheway, 2012). The fate of these byproducts is discussed in the following bullet.

- Residuals Management. In addition to the gas produced by the manufactured gas
 process, residual materials were also produced and separated from the gas at several
 steps during the process. These residuals were intermediate waste streams typically
 managed on-Site and further processed to create byproducts for disposal or reuse.
 Residuals from the manufactured gas process included the following:
 - Tar-Water Emulsion. Tar removed from the gas stream, particularly from the condenser, was often a tar-water emulsion. Tar required a low water content to be saleable. Tar-water emulsions were typically removed from clarification equipment and transported to residual management areas to separate the tar from the water (Hatheway, 2012). Tar and water were typically separated by placing the emulsion in pits, cisterns, or tar wells (typically shallow boxes that may be lined or unlined) and allowing the tar to settle out. A former plant map shows tar wells and a residue cistern located west of the purifiers near the edge of the ravine adjacent to the former gas works (Former Ravine). A former resident recalled a tar pit located on the southwest corner of the Former Gas Works Property (b) 2014), and an engineer's report (Tymstra, 1942) noted, "The tar emulsion is dumped in shallow pits dug at random in the ground." A historical journal (Perry, 2002) indicated that the former gas works "had a pond for dumping surplus creosote-type fluids. This would overflow and the material would go into the channel." It is unknown how tar-water emulsions were transported to these areas or how tar was transported from these areas to the tar storage tank, which was located on the south side of the Former Gas Works Property.
 - Storage, Distribution, and Disposal of Gas and Byproducts. Following purification, finished gas was stored and distributed via underground piping to the gas service area. Finished gas and byproducts of the manufactured gas process were collected, stored, and used or disposed of as follows:
 - Finished Gas. Gas that had passed through the scrubbers and purifiers was pumped through compressors located in the engine room (south of the main plant building) and stored in finished gas storage tanks located south of the main operations area. Gas was piped from the finished gas tanks to the gas distribution system along an 8-inch-diameter gas main located in Thompson Avenue. Typically, in manufactured gas distribution systems, a minor amount of oil would condense within the initial section of distribution piping, which would be collected in a drip tank located near the facility (Hatheway, 2012). A drip tank located just south of the Former Gas Works Property (Figure 2-3) is shown on a historical plant sketch.
 - Light Oil. Light oils typically contain one- or two-ring aromatic compounds, such as BTEX, and naphthalenes, and have a density less than

that of water (i.e., light, non-aqueous phase liquids [LNAPLs]). Light oils were sometimes reused in the carbureted water-gas process. According to a former worker, light oils were produced in small quantities at the former gas works and stored in a tank south of the finished gas storage tanks. The worker recalled lights oils were used as automotive fuel for workers' vehicles and were occasionally sprayed to control weeds in the southwest corner of the Former Gas Works Property (Simonson, 1997b).

- Carbureted Water-Gas Tar. This tar typically contains both light aromatics (e.g., BTEX) and semivolatile hydrocarbons. Semivolatiles in carbureted water-gas tar primarily consist of PAHs, but also include phenols and heterocyclic aromatics (i.e., carbazole or dibenzofuran). Carbureted water-gas tar is typically denser than water (i.e., dense non-aqueous phase liquids [DNAPLs]). According to a former worker (Simonson, 1997b), tar was a saleable product that was collected, stored in a tank on the south side of the Former Gas Works Property, and piped to barges at the Former Gas Works Dock. However, it is unlikely that all tar generated over the entire life span of the former gas works was recovered and sold in this manner, and some may have been accidentally spilled or released.
- Gas Liquor. Gas liquor is water containing dissolved and suspended tar and oil constituents. According to a 1942 report, gas liquor was reportedly discharged to the Port Washington Narrows through a drainpipe (Tymstra,1942), but it is unknown if this practice continued for any length of time.⁵
- Ash, Clinker, and Slag (Mineral Residue of Fuel and Feedstocks) from Furnaces. Ash is generally powdery, whereas clinker is partially fused, and slag is fused. Some of these materials were reportedly placed on the bluff along the shoreline north of the Former Gas Works Property

 (b) (6) 2014), and some may have been deposited in the Former Ravine.
- Soot from Furnaces. According to a 1942 report, this material was reportedly placed in the Former Ravine near the oil storage tanks (Tymstra, 1942), but it is unknown if this practice continued for any length of time.
- Spent Scrubber and Purifier Media. When scrubber and purifier media such as tar-soaked wood chips and shavings were saturated, they were removed and replaced. Spent scrubber media contains tar, and spent purifier media often contains tar, sulfide, and cyanide compounds removed during purification, including Prussian Blue (an iron-cyanide compound) (Hatheway, 2012). According to a 1942 report, tar-soaked wood chips and excelsior produced on-site were reportedly placed in the Former Ravine near the oil storage tanks (Tymstra, 1942), but it is unknown if this practice continued for any length of time. An individual who worked at the

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⁵ It is suspected that the drain pipe referred to in the 1942 report corresponds to the former outfall that was removed and plugged as part of the 2010 TCRA (see Section 3.3.1).

⁶ Boring logs for SP01 and MW04, which were located in the Former Ravine, indicate ash.

former gas works between 1953 and 1955 indicated that the spent purifier media were hauled off-Site (Simonson, 1997b).

2.3.1.2 Post-1972 Operations on the McConkey Property

Operations on the McConkey Property after the former gas works discontinued operations have included activities by Lent's between approximately 1972 and 1982, and industrial park operations by others from approximately 1982 to the present. Operations on the McConkey Property have included metal fabrication and sandblasting in the southern portion of the property, and parking and equipment storage across the other portion of the property. Two warehouse buildings are in the southern portion of the McConkey Property; the buildings are rented to separate tenants for storage of motor vehicles, vehicle parts, and associated mechanical equipment and tools. Historical and current operations on the McConkey Property are shown on Figure 2-4. A generalized process flow diagram of the metal fabrication process is shown on Figure 2-5.

Ecology inspected industrial park operations on the McConkey Property in 1992, 1993, 1994, and 1995, and observed the following activities during that period that may have resulted in contaminant releases:

- Improper storage of sandblast grit, solvents, and paint sludge at a metal-fabricating shop; and
- Debris and drums containing oily substances scattered around the industrial park.

2.3.1.3 Operations on the Sesko Property

The Sesko Property was used for bulk petroleum storage and distribution from as early as 1946 to no later than 1993, when the aboveground storage tanks (ASTs) were removed. Lent's was the primary operator of the tank farm on the Sesko Property. Former AST locations are shown on Figure 2-4. A process flow diagram of petroleum storage and distribution operations is provided on Figure 2-5. After 1993, the Sesko Property was used for boat maintenance, automobile salvage, equipment and debris storage, parking, and metal reclamation. The owner of the Sesko Property was involved in legal disputes with the City over nonconforming use of the Sesko Property (as a junkyard), violations of the Shoreline Management Act, and, in 2003, improper decommissioning of an underground storage tank (UST). Ecology spill records also indicate that approximately 25 gallons of gasoline were released to surface water from the Sesko Property in January 2003. Most of the equipment and debris has been removed, and the Sesko Property is currently vacant.

The Sesko Property includes remnants of the Former Ravine, which has been filled over the years. Fill activities have included the following:

• **Before 1930.** No records documenting fill activities before operation of the former gas works have been identified. However, based on a comparison of the 1919 shoreline (Figure 2-4) with an aerial photograph dated 1946 and sewer maps dated 1939, it appears that a portion of the Former Ravine was likely filled by the late 1930s, before

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⁷ Based on City directory information, Lent's continued operating on the McConkey Property for at least 3 years after the McConkeys acquired most of the McConkey Property in 1979.

construction of a historical residence located on the Sesko Property and before construction of the Lent's tank farm.

- 1931 to 1955. Aerial photographs and recorded observations (Tymstra, 1942; Judd, 2014) indicate that the western portion of the Former Ravine was filled between 1931 and 1955. Recorded observations indicate that people unaffiliated with the former gas works dumped miscellaneous garbage, trash, and fill in the Former Ravine before 1942. Residual materials from former gas works operations (i.e., soot, ashes, cinders, and tar-laden wood chips and shavings) were also reportedly dumped in the Former Ravine during this period (see Section 2.3.1.1).
- 1941 to 1974. An easement granted by Western to the City gave the City the right to dump refuse, garbage, and ashes from an incinerator into the Former Ravine. The easement reserved the right for Western to dump ashes and cinders in the easement area, which included the eastern 25 feet of the Former Gas Works Property (most of which lies on the current Sesko Property). According to the City, the historical records that partially document this time period were destroyed in a fire, and any documents regarding construction of the incinerator or dumping of refuse, garbage, or incinerator ash into the Former Ravine would have been lost in that fire.
- 1968. A DNR inspection reported that concrete and piping debris were placed in the Former Ravine (DNR, 1968).

Petroleum transfer lines that connected a dock located on the northern edge of the Sesko Property (Former Sesko Dock) to the Former ARCO Property and the Lent's tank farm were formerly located on the Sesko Property and may still be in place. An employee of the owner of the Sesko Property indicated that he had removed a portion of underground petroleum transfer piping he encountered in the northern portion of the Sesko Property. Petroleum transfer lines also reportedly connected the Former Sesko Dock to the Former SC Fuels Property to the east. Approximate pipeline locations, shown on Figure 2-4, were identified on construction plans for City sewer improvements (CH2MHill, 1982; MH&A, 1982).

2.3.2 Adjoining Properties

Surrounding properties include (1) the Penn Plaza Property, located to the south of the McConkey Property, (2) the Former ARCO Property, located to the west of the McConkey Property across Thompson Drive, and (3) the Former SC Fuels Property, located to the east of the Sesko Property across Pennsylvania Avenue (Figure 2-1). Historical and current operations on these properties are discussed in the following Sections.

2.3.2.1 Penn Plaza Property

There are five buildings on the Penn Plaza Property, which is used as an industrial park. Multiple tenants occupy the industrial park. Based on available records, the Penn Plaza Property has been used for commercial and/or industrial uses since the late 1930s or early 1940s. Prior to this time, an intermittent stream ran northeast across the Penn Plaza Property toward the Former Ravine on the current Sesko Property. This stream was reportedly used by area residents for dumping refuse and was filled in by 1942 (b) 2014).

Operations on the Penn Plaza Property have included Lent's operations from the 1940s to approximately 1985, and industrial park operations from approximately 1985 to the present. Lent's operations on the Penn Plaza Property included spray painting, metal plating, a pipe shop, truck repair, and parking for petroleum distribution. A former employee of Cascade, who worked in Bremerton in 1968 and 1969, recalled that wood treating may also have occurred as part of Lent's operations (Clapp, 1997). Since the cessation of Lent's operations, multiple tenants have used the Penn Plaza Property for industrial uses, including sheet metal fabrication, floating pier and acrylic septic tank manufacturing, concrete pipe/manhole manufacturing, heating and air conditioning repair, and marine propeller repair (TechLaw, 2006; Hart Crowser, 2007).

Ecology inspected operations at the Penn Plaza Property in 1992, 1993, 1994, and 1995, and identified the following activities that may have resulted in contaminant releases:

- A tenant reported to Ecology that an electroplating operation had made illegal discharges to a storm drain that resulted in a sewer backup.
- Ecology observed improper storage of waste concrete and waste oil at one of the tenant locations.
- Ecology observed diesel staining on the ground at another tenant location.
- Ecology observed debris and drums containing oily substances scattered around the industrial park.

On the north end of the Penn Plaza Property are oil and gasoline supply pipelines that connected the Former Sesko Dock with the Former ARCO Property to the west. The approximately location of these pipelines, based on a utility locate conducted during the time critical removal action (TCRA) in 2010, is shown on Figure 2-4.

2.3.2.2 Former ARCO Property

The Former ARCO Property was used for bulk petroleum storage and distribution from the mid-1940s to the late 1980s or early 1990s. Initially, 4 ASTs were present, with 2 added prior to 1956, 5 added in the late 1970s, and 4 added in the early 1980s for a total of 15 ASTs. Loading racks were located in the southeast corner of the Former ARCO Property. All tanks were removed by 1993. Property records indicate storage of gasoline, diesel, and oil. Product lines connected the ASTs on the Former ARCO Property with the Former Sesko Dock. Piping from the Former ARCO Property crossed the adjacent property to the north and ran west along the waterfront to a former dock (Former ARCO Dock) located approximately where the Port Washington Marina is today (see Section 2.3.3). According to a former resident, the piping to the Former ARCO Dock was located above ground (Judd 2014).

Since the early 1990s, the Former ARCO Property has been sporadically occupied by various tenants, including a tenant that conducted furniture refinishing and repair. The

⁸ Based on City directory information, Lent's continued operating on the McConkey Property for at least 3 years after the property was sold in 1979.

⁹ Petroleum for Lent's petroleum distribution was stored on what is now the Sesko Property.

Former ARCO Property is currently being used for commercial purposes by Pipeworks Mechanical and Service, Inc.

2.3.2.3 Former SC Fuels Property

The Former SC Fuels Property was used for bulk petroleum storage and distribution from the mid-1940s to the present. Operations on the Former SC Fuels Property are currently inactive. Initially, five ASTs were present, with one AST added prior to 1963, for a total of six ASTs. Four USTs were removed in 2003. Property records indicate storage of gasoline, diesel, and waste oil.

The Former SC Fuels Property is registered in Ecology's Voluntary Cleanup Program. A series of environmental investigations and remedial actions performed between 1997 and 2007 have confirmed releases of petroleum products and associated constituents, including gasoline, diesel, oil, BTEX, and PAHs. Additional information about the investigations and remedial actions is provided in Section 3.4.

Stormwater at the Former SC Fuels Property is collected in a series of catch basins, piped to an oil-water separator located at the top of the bluff, and discharged through an outfall to the Port Washington Narrows (Figure 2-4). Ecology conducted a Site visit in 2006, and noted a "gasoline odor" along the shoreline of the Former SC Fuels Property close to the stormwater outfall.

Pipes supplying petroleum to the Former SC Fuels Property tank farm ran from the Former SC Fuels Dock (see Section 2.3.3). An unknown number of petroleum transfer pipes also reportedly ran from the Former Sesko Dock to the tank farm on the Former SC Fuels Property, although their alignment is unknown (see Section 2.3.1.3).

2.3.3 Aquatic Parcels

Four docks were constructed in the aquatic parcels located adjacent (or closest to) to the properties described in Sections 2.3.1 and 2.3.2 (Figure 2-4). These aquatic parcels were leased from DNR. A description and brief history of each dock is included in the following Sections, and a detailed lease history prepared by DNR is provided in Appendix D

2.3.3.1 Former Gas Works Dock

The Former Gas Works Dock was constructed by Western on November 25, 1930, as part of the development of the former gas works. It was located on the aquatic parcel adjacent and to the north of the Former Gas Works Property. The Former Gas Works Dock was used to offload coal, briquettes, and oil (via a 3-inch-diameter pipeline). Records indicate that the Former Gas Works Dock was also used to transfer heavy-end byproducts. In 1948, as part of the propane blending retrofit, the Former Gas Works Dock was updated to allow offloading of propane gas. Based on review of aerial photography, the Former Gas Works Dock was removed sometime between 1971 and 1974.

2.3.3.2 Former ARCO Dock

The Former ARCO Dock was constructed by the Richfield Oil Corporation in approximately 1942. It was located on the aquatic parcel immediately adjacent and to the

west of the aquatic parcel operated by the former gas works. The Former ARCO Dock served as both boat moorage and support for the pipelines associated with upland ARCO operations. It was removed by Richfield Oil's successor in the mid-1980s.

2.3.3.3 Former Sesko Dock

The Former Sesko Dock was constructed by Lent's in approximately 1942. It was located on the aquatic parcel immediately adjacent and to the east of the aquatic parcel operated by the former gas works. The Former Sesko Dock was used to support supply pipelines for barge delivery of diesel and stove oil, which were stored on the Sesko Property. During the 1970s and 1980s, the Former Sesko Dock was also used to supply the tank farm on the Former ARCO Property and the tank farm on the Former SC Fuels Property. In 1993, the pipelines on the Former Sesko Dock were removed. The Former Sesko Dock was removed in September 2001 pursuant to a DNR order.

2.3.3.4 Former SC Fuels Dock

The Former SC Fuels Dock was constructed by General Petroleum Corporation of California in 1942. It was located on the aquatic parcel immediately adjacent and to the east of the aquatic parcel where the Former Sesko Dock was located. The Former SC Fuels Dock was constructed for handling petroleum products. The Former SC Fuels Dock was removed in 1967 by Mobil Oil Corporation when barge deliveries of petroleum products were discontinued.

2.4 Environmental Setting

2.4.1 Climate and Meteorology

The Bremerton, Washington, area is dominated by a marine temperate climate with cool and comparatively dry summers and mild, wet, and cloudy winters (WRCC, 2014). The average annual high temperature for Bremerton is 60 degrees Fahrenheit (°F), and the average annual low temperature is 43°F (WRCC, 2014). Average annual precipitation is 52 inches, with nearly half of that occurring in November, December, and January (WRCC, 2014). During this wet season, rainfall is usually light to moderate in intensity and continuous over a period of time, rather than brief, heavy downpours. During the driest months of July and August, it is not unusual for 2 to 4 weeks to pass with only a few showers (WRCC, 2014). The prevailing wind direction in the region is south or southwest during the wet season and northwest in summer, with an average wind velocity of less than 10 miles per hour (WRCC, 2014).

2.4.2 Topography and Drainage

The Former Gas Works Property is located on a bluff on the south shore of the Port Washington Narrows. The Former Gas Works Property generally slopes gently to the north and is covered with buildings or pavement. At the northern edge of the Former Gas Works Property, a vegetated bluff slopes steeply down to the beach. Over time, the bluff has expanded to the north with the placement of fill material. Remains of the Former Ravine along the eastern edge of the Former Gas Works Property can be seen as a cove located at the northern edge of the Sesko Property. Stormwater drainage characteristics on the Former Gas Works Property and adjacent properties are as follows:

- McConkey and Penn Plaza Properties. Pavement covers most of the McConkey and Penn Plaza Properties, and the properties have catch basins connected to the City stormwater drainage system. A City stormwater and combined sewer overflow (CSO) outfall is located offshore, north of Pennsylvania Avenue. A catch basin in the northwest corner of the McConkey Property is connected to an outfall on the beach below the bluff.
- **Sesko Property.** Most of the Sesko Property is unpaved. Stormwater either infiltrates or runs off, presumably to the north toward the Port Washington Narrows.

2.5 Geology and Hydrogeology

2.5.1 Regional Geologic Setting

The Site lies within the Puget Lowland, an area that has alternated between glacial and interglacial environments during the last 2 million years. The result has been a stacked and imperfectly preserved sequence of glacial and nonglacial strata (Armstrong et al., 1965; Blunt et al., 1987; Booth et al., 2004a). This irregular stratification has been further impacted by the tectonics of the Seattle fault, a regional thrust fault system that extends through the area (Bucknam et al., 1992; Johnson et al, 1999; Blakely et al., 2002), including a strand through Oyster Bay (Washington DNR, 2014). The impacts of the fault system include uplift and tilting of bedrock and Quaternary strata in some areas and subsidence in others (Nelson et al., 2003; Kelsey et al., 2004).

Interglacial climates produced sediments much like the forested Puget Lowland before extensive development, with broad floodplains and gently sloping uplands. These deposits include silty to sandy floodplain sediments, scattered gravelly channel deposits, and peat and lacustrine (lake) sediments. Glacial climates resulted in rapid accumulation of glacial sediments and scour of preexisting landforms and deposits. These deposits include advance glacial lake (glaciolacustrine) deposits, advance outwash (glacial river deposits), glacial till (subglacial deposits), and recessional glacial deposits.

Bedrock crops out on the northern end of the peninsulas between Phinney Bay and Ostrich Bay, and elsewhere generally north and west of the Site (Washington DNR, 2014). Map data and limited deep well data suggest that bedrock generally dips to the south and west below the Site area (Eungard, 2014). This bedrock dip forms a regional basement aquitard (Jones, 1998). Some of the older sediments above bedrock are also likely tipped in this direction due to regional rotation along the Seattle fault (Booth et al., 2004b). Younger deposits, including those encountered in explorations for this project, are expected to be generally more horizontal but will include a number of discontinuous and irregularly shaped lenses of fine- and coarse-grained sediments that will impact the velocity and direction of groundwater flow. A conceptual geologic model of the Site area, including surficial geology (Figure 2-6) and subsurface geology (Cross Section AA–AA' on Figure 2-7) has been developed using regional map and well log data. Areas below the known exploration depths are shown as "undifferentiated."

The conceptual regional hydrogeologic model is one of rainfall and infiltration on an upland covered generally with till and glacial outwash. Some of this water runs off as stormwater, while a portion infiltrates. The water that infiltrates (groundwater) will migrate more quickly through more-permeable strata and will be generally retarded by

less-permeable strata. The migration of water through these strata is influenced by the location and dip of the low-permeability strata (aquitards), as well as the location of waterways and other low-lying areas, which are often points of groundwater discharge. Regional patterns indicate that uplands are generally recharge areas, and slopes near sea level are discharge points. Groundwater also migrates from deeper strata and discharge upward into waterways.

2.5.2 Site Geology

Four principal geologic units have been identified based on previous explorations: fill, natural glacial deposits of the Vashon Drift, nonglacial deposits from one or more of the interglacial events that preceded the Vashon glaciation, and deposits from an older glaciation. The characteristics and distribution of these major sequences are described in this Section, from the stratigraphic top (generally younger) to the bottom. Note that these geologic interpretations are based on logs prepared by multiple geologists over the course of the prior investigations. Subsurface interpretations from these earlier explorations (e.g., fill characteristics or extent) may be refined later based on future observations.

The locations of the cross sections are shown on Figure 2-8, and four geologic cross sections are provided on Figures 2-9 through 2-12. Soil boring logs are provided in Appendix E. A description of the soils observed at the Site is provided in the following text.

Although fill was not specifically identified in many of the soil boring logs, it was apparently present in the majority of the previous explorations at the Site, in thicknesses ranging from a foot or less to about 15 feet. The thickest fill is present in the Former Ravine area on the Sesko Property. Fill is generally composed of brown to black, loose to very dense, or stiff to very stiff variable mixtures of silt and sand with variable amounts of gravel, coal fragments, asphaltic concrete, and other debris. The density and consistency of the fill was generally high for nonstructurally placed fills and may be due to inclusion of ash in the fill soils, which can produce slight cementation of soils.

Over most of the Site, glacial deposits were encountered beneath the surficial fill. The geologic maps of the Site indicate the glacial unit is the Vashon Drift. The soils encountered in the explorations generally consisted of clean (fines are absent) to silty fine-to medium-grained sand with trace to minor amounts of gravel and scattered interbeds of sandy silt. These glacial deposits were observed to be dense to very dense and were generally brown to gray. The gradation and density of this unit suggests that it is primarily Vashon advance glacial outwash. This unit has moderate permeability and, where saturated, will form an aquifer. The thickness of this unit at the Site ranges from 10 to 35 feet.

Pre-Fraser nonglacial deposits (predating the Vashon Glaciation) are present in the bluffs and uplands in the northeastern portion of the Site. Explorations encountered olive to gray and brown, stiff to hard silt to sandy silt with interbeds of very dense silty sand ranging in thickness from 2 to 10 feet. Thin interbeds or lenses of clay and silty clay and scattered gravelly layers may be present. This unit generally has low permeability; however, cleaner sandy layers may become saturated.

An older glacial sequence is present below the Vashon outwash and the pre-Fraser nonglacial deposits. The thickness of this unit has not been defined at the Site. The older glacial sequence consists of lenses or discontinuous layers of glacial till within an outwash-like brown to gray, very dense slightly silty to silty sand. The lenses of till are composed of brown to gray very dense silty gravel with sand and silty sand with gravel. The till lenses are generally considered an aquitard, but the outwash-like silty sand component was noted to be wet below about the 5- to 10-foot elevation, which probably reflects the regional water table. The scope of work for the RI, as described herein, will include additional investigations to determine whether the till acts as an aquitard at the Site.

2.5.3 Hydrogeology

Groundwater on the McConkey Property and Sesko Property was encountered at depths between 15 and 41 feet. Groundwater elevations have ranged between 3 and 10 feet above mean sea level, with an estimated flow direction to the north-northwest (to the Port Washington Narrows) during one sampling event (GeoEngineers, 2007b). Monitoring well construction details and groundwater elevation measurements are summarized in Table 2-1. Well construction logs are included in Appendix E.

Groundwater on the Former SC Fuels Property has been encountered at depths between 4 and 15 feet, with an estimated flow direction to the northwest. Groundwater on the Former SC Fuels Property appears to be perched within sandy zones present in generally low-permeability nonglacial soils.

The estimated directions of groundwater flow on the McConkey, Sesko, and Former SC Fuels Properties, based on previous studies, are shown on Figure 2-13. However, groundwater studies to date have not evaluated the effect of tidal influence on Site groundwater levels and flow direction. One-time groundwater elevation measurements are prone to error if tidal effects are significant.

2.6 Human Populations and Land Use

The Former Gas Works Property is in Bremerton, which is the largest city on the Kitsap Peninsula and home to Puget Sound Naval Shipyard and the Bremerton Annex of Naval Kitsap Base. According to the 2010 census, the population of Bremerton is 37,729 people with 1,328 inhabitants per square mile. The racial makeup of Bremerton is predominantly white/Caucasian (74%) with the rest of the population classified as "other" or two or more races (10.4%), African American (6.7%), Asian (5.5%), Native American (2.0%), and Pacific Islander (1.3%). According to the Tribe government website, the total population of the Tribe is 950 people.

The Former Gas Works Property is in an area of industrial-zoned properties that includes the Former ARCO Property and Former SC Fuels Property. Surrounding this industrial property core are residential properties and a marina. A zoning map is included on Figure-2-1. The Former Gas Works Property is immediately adjacent to intertidal sediments and surface water within the Port Washington Narrows, which can be accessed by the public.

2.6.1 Tribal Use

Tribal commercial, subsistence, and ceremonial fisheries have historically occurred in Dyes Inlet and the Port Washington Narrows. The Tribe has stated that "Suquamish tribal members fully intend to continue to fish these areas for cultural, subsistence and commercial purposes" (Suquamish Tribe, 2014). According to the Tribe, it "uses the Washington Commercial Shellfish Growing Area Classification to determine the suitability of bivalve harvests (i.e., claims, oysters)" (Suquamish, 2011). The marine area adjacent to the Former Gas Works Property is designated as "Unclassified" due to the proximity of CSOs, which precludes shellfish harvesting. However, according to the Tribe, the harvest of finfish and other marine invertebrates (i.e., crab and sea cucumber) are not restricted adjacent to the Former Gas Works Property (Suquamish, 2011).

2.6.2 Drinking Water Use

Water services at the Site and surrounding area are supplied by the City. The closest public water supply wells are located over one mile from the Site. The use of private wells within the Bremerton Water Service Area is not allowed, and there are no drinking water wells near the Site listed in Ecology's database.

The Site is located adjacent to the Port Washington Narrows, a saltwater body. The extent of saltwater intrusion and the potability of Site groundwater, and its potential future use as a drinking water source will be evaluated as part of the RI.

2.7 Port Washington Narrows and Dyes Inlet

The Former Gas Works Property is located along the Port Washington Narrows, which is a tidal channel connecting Dyes Inlet to Sinclair Inlet and Puget Sound. Dyes Inlet is a terminal estuary, comprising five embayments (Phinney, Mud, Ostrich, Oyster, and Chico Bays) and the Port Washington Narrows (Figure 2-14).

The waters of Port Washington Narrows are relatively shallow, with average depths of less than 30 feet. Depths within Dyes Inlet range up to 100 feet, but are typically less than 50 feet. Area bathymetry is shown on Figure 2-14.

The shorelines of the Port Washington Narrows and Dyes Inlet have been extensively developed. These shorelines include the cities of Bremerton and Silverdale as well as the community of Tracyton. Other significant features include several former U.S. Navy facilities and regional transportation networks, including State Routes 3 and 303. The Warren Avenue and Manette Bridges are located across the Port Washington Narrows east of the Former Gas Works Property.

Hydrologic inputs to the Port Washington Narrows and Dyes Inlet include the tidal exchange with Sinclair Inlet and freshwater inflows from both stream and piped flows. Information from Kitsap County and the City regarding identified stormwater outfalls, CSO discharge points, and surface water inputs is summarized on Figure 2-14. Additional private and municipal outfalls may be present in addition to those identified by these information sources.

Hydraulic exchange between Dyes Inlet, the Port Washington Narrows, and the balance of Puget Sound is limited by the geography and the resulting hydrodynamics. In addition to

tide and current data available from public sources (e.g., National Oceanographic and Atmospheric Administration [NOAA]), the waters of Dyes Inlet and the Port Washington Narrows have been studied as part of regional water quality programs. Total maximum daily load (TMDL) studies and a contaminant mass balance evaluation have been performed for Dyes Inlet and may provide useful data for the RI/FS. Hydrodynamic modeling of the area has been performed as part of regional studies of Puget Sound. The results of additional studies are available to characterize environmental quality within Sinclair Inlet, immediately south of Dyes Inlet and the Port Washington Narrows. The Sinclair Inlet studies include extensive testing that has been performed in association with the Bremerton Naval Shipyard, as well as other regional study programs. These studies and evaluations are further addressed in Sections 3.5 and 3.9.

2.8 Natural Resources

This Section describes the natural resources of the upland areas, aquatic habitats, and related data needs for the RI/FS.

2.8.1 Upland Areas

The upland areas of the Former Gas Works Property and surrounding areas have been developed for industrial uses consistent with zoning provisions. However, some terrestrial and riparian habitat is present, particularly on the bank adjacent to the Port Washington Narrows, the Former Ravine, and the shoreline areas of the McConkey and Sesko Properties. The Washington Department of Fish and Wildlife (WDFW) manages a Priority Habitats and Species Program (PHS). Preliminary queries of WDFW's PHS system did not identify any priority terrestrial natural resources on the parcels associated with the Former Gas Works Property

2.8.2 Aquatic Habitats

Aquatic habitats at the Site include those in the beach and subtidal areas within and near the Former Gas Works Property. Shoreline and aquatic habitat adjacent to the Former Gas Works Property are located within the Tribe's Usual and Accustomed area. Fish and shellfish resources are present within the waters of the Port Washington Narrows and Dyes Inlet. Fish and crab are known to be present and support commercial, recreational, and tribal fisheries. Shellfish harvesting within the Port Washington Narrows and Dyes Inlet has been restricted due to water-quality-related shellfish harvesting closures. However, efforts have been made by state and local governments, tribes, and other stakeholders to improve water quality in the area and reduce or lift these shellfish harvesting restrictions. A number of shellfish enhancement projects have been proposed within portions of Dyes Inlet. It is not known what measures have been undertaken by the Washington State Department of Health (WDOH) or the Kitsap Public Health District (KPHD) to monitor illicit shellfish harvesting within Dyes Inlet or the intertidal areas adjacent to the Site. Signage indicating the closure of the beach adjacent to the Former Gas Works Property was installed as part of the 2013 TCRA (see Section 3.3.2).

The query of the WDFW PHS identified two aquatic natural resources in the vicinity of the Former Gas Works Property: estuarine intertidal aquatic habitat along the northern and southern shorelines of the Port Washington Narrows and hardshell clams along the northern shoreline of the Port Washington Narrows.

2.9 Cultural Resources

There are no recorded archaeological sites or historic structures at the Former Gas Works Property or in the immediate vicinity. However, no cultural resources surveys have been conducted on the Site or in the vicinity prior to the present project. The documented archaeological sites nearest to the Former Gas Works Property include the following:

- Site 45KP121, a precontact and historic-era shell midden site, is located in Evergreen Park, approximately 0.6 miles east-southeast of the project area;
- The Manette Site (45KP009), a large precontact midden and possible fortification site where human remains have reportedly been found, is located on a bluff above the beach, just west of the Manette Bridge (1.2 miles east-southeast of the project area); and
- A number of ethnographic place names have been recorded at various locations along the Port Washington Narrows.

Kitsap County assessor's records (accessed January 2014) indicate that there is one building older than 50 years on the Penn Plaza Property—a warehouse constructed in 1955. The structure has not been evaluated for National Register of Historic Places (NRHP) eligibility. No impacts on this structure are anticipated during the RI/FS.

An archaeologist from Anchor QEA, LLC (Anchor) visited the project area in August 2013 to make a preliminary assessment of current conditions. The project area has been extensively modified in the historic and modern eras, with placement of fill materials and debris, and development and redevelopment of the Site for industrial uses. No native sediments, other than active beach deposits, were visible in the project area.

3 Initial Evaluation

This Section summarizes the regulatory requirements and existing data that supported the development of the preliminary conceptual site model (CSM), which is described in detail in Section 4.

3.1 Regulatory Requirements

This Section identifies initial applicable or relevant and appropriate requirements (ARARs), preliminary remediation goals (PRGs), and remedial action objectives (RAOs) for the purposes of project planning. Potential ARARs were identified to facilitate communications with support agencies, help plan potential field activities, and assist in the identification of RAOs and PRGs. Initial PRGs were identified to help evaluate existing data and assist in the selection of appropriate analytical methods. The ARARs, PRGs, and RAOs will be further developed during the RI/FS process. Those ARARs, PRGs, and RAOs that are determined to be applicable to the Site-related decisions may include some, none, or all of those identified in this Section. The ARARs, PRGs, and RAOs that are ultimately determined to be applicable to the Site-related decisions will be established in consultation and coordination with key stakeholders and the public during the RI/FS process.

3.1.1 Applicable or Relevant and Appropriate Requirements

The project must comply with CERCLA Section 121, which requires remedial actions to achieve ARARs. According to the National Contingency Plan (Code of Federal Regulations, Title 40, Section 300.5 [40 CFR 300.5]), applicable requirements are those cleanup standards, standards of control, and other substantive requirements, criteria, or limitations promulgated under federal environmental or state environmental and facility siting laws that specifically address a hazardous substance, pollutant, contaminant, remedial action, location, or other circumstance identified at a CERCLA site. Appropriate and relevant requirements are cleanup standards, standards of control, and other substantive requirements, criteria, or limitations promulgated under federal environmental or state environmental or facility siting laws that are not applicable to a hazardous substance, pollutant, contaminant, remedial action, location or other circumstances at a CERCLA site, but address problems or situations similar to those encountered at the site that their use is well suited to the particular CERCLA site.

Some federal, state, and local environmental and health agencies may develop criteria, advisories, guidance documents, and proposed standards that are not legally enforceable but contain useful information for selecting cleanup levels or implementing a cleanup remedy. These fall into the category of "to be considered" (TBC) elements. TBCs are not mandatory requirements but may complement the identified ARARs.

ARARs and TBCs potentially relevant to the RI/FS are presented in Tables 3-1 through 3-3 and organized into the following categories:

- Contaminant-specific requirements;
- Location-specific requirements; and

• Performance, design, or other action-specific requirements.

Some ARARs fit neatly into a single category, whereas others may fall into more than one category. The categories are described as follows:

- Contaminant-specific ARARs are laws and requirements that establish health- or risk-based numerical values or methodologies for developing such values (EPA, 1988b).
 These ARARs are used to establish the acceptable concentration of a contaminant that may remain in or be discharged to the environment. As such, contaminant-specific ARARs are considered in identifying the PRGs. Contaminant-specific ARARs are listed in Table 3-1.
- Location-specific ARARs are requirements that are triggered based on the location of
 the remedial action to be undertaken (EPA, 1988b). Location-specific ARARs may
 restrict or preclude certain remedial actions or may apply only to certain portions of
 the Site. Some location-specific ARARs overlap action-specific ARARs. Locationspecific ARARs are listed in Table 3-2. An example of a location-specific ARAR is
 the Point Elliott Treaty of 1855.
- Action-specific ARARs are performance, design, or other requirements that may place
 controls or restrictions on a remedial action (EPA, 1988b). Action-specific ARARs are
 typically technology- or activity-based requirements or limitations on actions, and
 these requirements may include contaminant-specific standards or criteria that must be
 met as the result of an action. For remedial actions at the Site, these requirements are
 not necessarily triggered by the presence of specific contaminants in Site media, but
 rather by the specific actions that occur at the Site. Action-specific ARARs are listed
 in Table 3-3.

3.1.2 Remedial Action Objectives

RAOs consist of goals for protecting human health and the environment that are specific for each potentially contaminated environmental medium (e.g., soil, groundwater, and sediment). RAOs for protection of human receptors typically include both a contaminant level and an exposure route. RAOs for protection of environmental receptors typically seek to preserve or restore a resource and are typically expressed in terms of the medium of interest and target cleanup levels. The preliminary RAOs related to the protection of human health are as follows:

- **Groundwater.** Reduce risk to human health from direct contact with, and consumption of, groundwater contaminated with Site-related contaminants of concern (COCs)¹⁰ to protective levels.
- **Sediment.** Reduce risk to human health from consumption of fish and shellfish containing Site-related COCs to protective levels.
- **Sediment.** Reduce to risk to human health from incidental ingestion and/or dermal exposure to Site-related COCs during potential recreational use of the beach areas at the Site to protective levels.

¹⁰ Under CERCLA guidance, those COPCs identified as posting unacceptable risk during the baseline risk assessment should be retained as contaminants of concern (COCs) for further evaluation of remedial options during the FS stage of the RI/FS.

- **Vapor.** Reduce risk to human health from inhalation of vapors from groundwater and/or soils contaminated with Site-related COCs to protective levels.
- Soils (Surface and Subsurface). Reduce risk to human health from direct contact with or incidental ingestion of Site-related COCs to protective levels.

The preliminary RAOs related to environmental protection are as follows:

- **Groundwater.** Reduce, to protective levels, risks to ecological receptors from direct contact with and consumption of groundwater contaminated with Site-related COCs that discharges to surface water at the shoreline, including indirect exposure from consumption of prey exposed to groundwater discharging to surface water.
- **Groundwater**. Reduce, to protective levels, migration of contaminants in groundwater to surface water or sediments.
- Upland Soil. Reduce, to protective levels, risks to terrestrial wildlife exposed to Siterelated COCs through direct contact with and incidental ingestion of Site soil or consumption of soil-dwelling invertebrates.
- **Sediment.** Reduce, to protective levels, risks to aquatic wildlife from exposure to Siterelated COCs in surface sediments or in prey species at the Site.
- **Sediment.** Reduce, to protective levels, risks to the benthos from Site-related COCs in surface sediments.

The preliminary RAOs will be developed further throughout the RI/FS process, in consultation with key stakeholders and the public, and may be revised, refined, or replaced.

3.1.3 Preliminary Remediation Goals

This Section identifies PRGs for the initial screening of existing soil, groundwater, and sediment data. The purpose for PRGs is to establish a preliminary screening level, based on ARARs, to evaluate investigation results. PRGs can be updated throughout the RI/FS process and any PRG screening does not affect the identification of COCs. Surface water initial PRGs have been identified to assist in the development of this Work Plan; however, no surface water data are available for the Site. The initial PRGs were used in the development of the SQAPPs (Appendices A and B) to select appropriate analytical methods.

Potential PRGs include numerical values identified in ARARs, peer-reviewed risk-based values, or values identified in other screening benchmark sources. Potential PRGs include values from the following sources:

1. ARARs:

- Soil: none available (except for those related to polychlorinated biphenyls (PCBs) in the Toxic Substances Control Act);
- Groundwater: maximum contaminant levels (MCLs);
- Surface water: Washington State-specific and EPA human health criteria (organisms only) promulgated under Section 303(c) of the Clean Water Act (CWA) (EPA, 2016)

and national recommended water quality criteria for human health (organisms only) and -aquatic life (acute and chronic)¹¹; and

• Sediment: Washington State Sediment Management Standards (SMS).

2. Peer-reviewed sources:

- Soil: EPA human health regional screening levels (RSLs) and EPA ecological soil screening levels (EcoSSLs);
- Groundwater: EPA human health RSLs;
- Surface water: none available; and
- Sediment: NOAA effect range-low (ER-L) and effect-range-medium (ER-M) benchmarks (Long et al., 1995).

3. Other screening benchmark sources:

- Soil: EPA Region 5 Resource Conservation and Recovery Act (RCRA) EcoSSLs
- Groundwater: none available;
- Surface water: EPA Region 3 Biological Technical Assistance Group (BTAG)
 ecological surface water screening benchmarks and EPA Region 5 RCRA ecological
 surface water screening levels; and
- Sediment: EPA Region 3 BTAG ecological sediment screening benchmarks and EPA Region 5 RCRA ecological sediment screening levels.

Tables 3-4, 3-5, 3-6, and 3-7 summarize the potential PRGs from these sources for each medium (soil, groundwater, sediment, and surface water, respectively) and identify an initial PRG for each contaminant. The initial PRG for a given contaminant was selected as the lowest of the ARARs or peer-reviewed risk-based criteria. If a value from these first two sources is unavailable, the initial PRG was selected as the lowest value in the "other screening benchmark" category. For sediment, the regionally specific SMS value was used. If no SMS value exists for the contaminant, the peer-reviewed NOAA value was used.

For soil, two different initial PRGs were identified: one for surface soil to a depth of 0- to -3 feet below ground surface (which includes a consideration of screening levels for terrestrial ecological receptors from 0- to 1-foot, and upland construction worker exposure scenario of 0- to 3-feet) and one for subsurface soil at depths below potential ecological exposures and construction worker scenarios. The initial PRGs include the following:

• Soil:

o EPA RSLs – residential

o EPA RSLs – industrial

o EPA EcoSSLs – birds

¹¹ The National Toxics Rule (NTR) 40 CFR131.36 establishes chemical-specific numeric criteria for priority toxic pollutants for certain states. Washington has been withdrawn from the NTR for those state-adopted criteria approved by EPA (EPA, 2016) but the NTR remains an ARAR for other criteria. However, for this project there are no compounds with NTR criteria that do not have state-adopted criteria so the NTR criteria are not included.

- o EPA EcoSSLs mammals
- o EPA EcoSSLs invertebrates
- o EPA EcoSSLs plants
- o EPA Region 5 RCRA EcoSSLs

Groundwater:

- o EPA MCLs
- EPA RSLs tap water

Sediment:

- Washington State SMS sediment cleanup objectives (SCOs);
- o NOAA ER-L benchmarks (Long et al., 1995);
- EPA Region 3 BTAG ecological marine sediment screening benchmarks;
 and
- o EPA Region 5 RCRA ecological sediment screening benchmarks.

Surface water:

- National recommended water quality criteria for human health (consumption only) and aquatic life (EPA, 2013b);
- EPA Region 3 BTAG ecological marine surface water screening benchmarks;
- o EPA Region 5 RCRA ecological surface water screening levels; and
- o CWA-Effective criteria for the protection of human health (consumption of organisms) (EPA, 2016).

4. Seafood Consumption-based PRGs:

Additional PRGs based on seafood consumption will be developed following further consultation with EPA and the Suquamish Tribe. The PRGs will consider potential risks to tribal members who consume fish or shellfish from the Site. The PRGs for these exposure scenarios will be developed in the Risk Assessment Technical Memorandum (see Section 5.3.1) and the Baseline Human Health Risk Assessment (Section 5.3.2). Additional testing beyond what is included in this Work Plan may be required to support development of these PRGs (e.g., sampling and analysis of seafood tissue). Where necessary, the scope of supplemental testing will be defined in the Risk Assessment Technical Memorandum.

3.2 Previous Site Investigations

Previous environmental field investigations at the Former Gas Works Property include the following:

- Sesko Property Field Inspection (Ecology, 1995);
- Preliminary Upland Assessment, McConkey and Sesko Properties (GeoEngineers, 2007b); and

• Targeted Brownfields Assessment (TBA), McConkey and Sesko Properties (E&E, 2009).

The upland exploration locations and sampling depths by analyte group are provided on Figure 3-1. The scope and general conclusions of each study are described in the following Sections.

3.2.1 Ecology Field Inspection (1995)

In 1995, Ecology collected three surface soil samples from the Sesko Property and one surface sediment sample from the tidelands just north of the Sesko Property. The samples were analyzed for metals and SVOCs. High concentrations of PAHs were detected. Ecology used the data in conducting a Site Hazard Assessment and gave the Site a ranking of "1" (highest concern).

3.2.2 Preliminary Upland Assessment (2007)

In 2007, on behalf of the City and funded by a brownfield grant from EPA, GeoEngineers conducted a preliminary assessment of the McConkey and Sesko Properties (GeoEngineers 2007a) that included the following:

- Advancing eight soil borings and collecting soil samples to a maximum depth of 45 feet:
- Installing monitoring wells at each of the eight soil boring locations and collecting groundwater samples; and
- Analyzing soil and groundwater samples for total petroleum hydrocarbons (TPH), volatile organic compounds (VOCs), SVOCs, PCBs, and metals.

This work identified relatively high concentrations of gasoline- and diesel-range TPH, VOCs including benzene, and PAHs in soil and groundwater on the McConkey and Sesko Properties. VOCs and PAHs were detected in soil samples at depths up to 35 feet. Several metals, including arsenic, lead, and chromium (including chromium VI), were detected in groundwater at concentrations greater than the potential drinking water cleanup standards.

3.2.3 Targeted Brownfield Assessment (2008)

In 2008, on behalf of EPA, Ecology & Environment, Inc. (E&E) conducted a TBA of the McConkey and Sesko Properties (E&E, 2008) that included the following:

- Advancing seven soil borings and collecting soil samples to a maximum depth of 45 feet:
- Installing monitoring wells at two of the seven boring locations:
- Collecting groundwater samples from the two wells and from temporary screens placed at four of the seven soil boring locations;
- Collecting five surface sediment samples from the beach north of the properties;

 Analyzing soil, groundwater, and sediment samples for TPH, VOCs, SVOCs, and metals.

Similar to the Preliminary Upland Assessment, this work identified relatively high concentrations of gasoline- and diesel-range TPH, VOCs including benzene, and PAHs in soil and groundwater on the McConkey and Sesko Properties. The assessment also identified relatively high concentrations of PAHs in surface sediments. VOCs and PAHs were detected in soil samples at depths up to 45 feet.

3.3 Previous Site Removal Actions

Two TCRAs have been performed at the Site as described in this Section.

3.3.1 Time Critical Removal Action (2010)

In August 2010, sheens on the surface water of the Port Washington Narrows were reported to KPHD. Upon further investigation, KPHD identified a 12-inch-diameter concrete pipe that appeared to be the source of the sheen. The pipe is believed to be an abandoned City CSO outfall. KPHD reported the release to EPA, which in turn notified the U.S. Coast Guard (USCG) for a response because the pipe was within its jurisdiction. In 2010, at the request of EPA, E&E conducted sampling and analysis as part of the EPA and USCG's initial response. The response sampling included the collection of 32 surface sediment samples from a depth of 0 to 6 inches. The sediment samples were analyzed for VOCs and SVOCs, both of which were detected.

EPA, DNR, KPHD, and Ecology entered into a USCG-led coordinated response under a Unified Command Structure. Cascade became aware of the response in October of 2010 and informed the USCG that it was interested in contributing to the response. USCG subsequently added Cascade to the Unified Command Structure and issued Cascade an Administrative Order for a Pollution Incident (Order) to implement response actions at the Site under the oversight of USCG. Cascade accepted the Order in a letter dated October 29, 2010.

In response to the Order, Cascade developed an Incident Action and TCRA Work Plan (Anchor QEA and Aspect 2010), which outlined the scope and details of the 2010 TCRA. The 2010 TCRA included the following key elements:

- Investigation of the location and orientation of the abandoned pipe;
- Permanent plugging of the pipe as close as practicable to the shoreline;
- Removal of all portions of the pipe from the new plug to the terminus of the pipe;
- Backfilling of the excavation created by removal of the pipe with clean beach material;
- Placement of an organoclay mat over impacted sediments (with minimal disturbance) near the terminus of the pipe that were observed to generate sheen; and
- Continued maintenance of a containment system until field observations and inspections confirm that the situation is stable (no sheen).

On November 5, 2010, USCG and the other members of the Unified Command Structure approved the Incident Action and TCRA Work Plan. Cascade commenced the 2010 TCRA immediately upon approval and completed the 2010 TCRA on November 8, 2010 (Anchor QEA, 2011). The removal action satisfied the following objectives of the Incident Action and TCRA Work Plan:

- The pipe was located and traced to the shoreline.
- The pipe was plugged as close as practicable to the shoreline, at the location specified in the Incident Action and TCRA Work Plan.
- All pipe sections downgradient of the new plug were removed together with all overburden sediments.
- All excavations were filled to grade with clean beach material.
- The organoclay mat was placed over the area of impacted sediments specified in the Incident Action and TCRA Work Plan.

Inspections of the 2010 TCRA area were completed as specified in the Incident Action and TCRA Work Plan. The inspections consisted of visual observation of the ground surface at the pipe plug area and the overlying organoclay mat to identify any potential surface sheen. Following completion of the removal action, inspections were performed at decreasing intervals over time, starting at biweekly intervals and decreasing to quarterly intervals. Inspections have continued on a quarterly basis. No surficial sheens related to the 2010 TCRA have been observed to date. The constructed elements of the 2010 TCRA are shown on Figure 3-2.

3.3.2 Time Critical Removal Action (2013)

In 2013, Cascade completed a removal evaluation pursuant to the requirements of the AOC and the EPA-approved Removal Evaluation Work Plan (Anchor QEA and Aspect, 2013a). The objective of the removal evaluation was to assess whether suspected migration pathways at the Site pose a threat to human health, welfare, or the environment if left unaddressed before completion of the RI/FS. The results of the removal evaluation were reported in the EPA-approved Removal Evaluation Report (Anchor QEA and Aspect, 2013c). The removal evaluation identified the following conditions that warranted action before completion of the RI/FS:

- Stormwater intrusion into Manhole A. Manhole A was believed to remain connected to the 12-inch-diameter concrete pipe that was plugged as part of the 2010 TCRA. Based on inspections conducted as part of the removal evaluation, it was determined that stormwater could have been entering Manhole A through surface runoff or via a piping connection to Manhole A from a nearby sump. Stormwater entering Manhole A posed a risk of hydraulically surcharging the pipe plugged during the 2010 TCRA, which in turn could have increased the risk of a hazardous substances release to the Port Washington Narrows.
- Hydrocarbon sheen and deposits of solid hydrocarbon material in SG-04/SG-05 Area. Hydrocarbon sheens were observed in shallow subsurface sediments in the western area of the beach, near sampling stations SG-04 and SG-05. Surficial solid hydrocarbon material was also observed in the SG-04/SG-05 area. Both the sediments

containing hydrocarbon sheen and the solid hydrocarbon material contained concentrations of PAH compounds that were elevated in comparison to those of the surrounding beach sediments.

The Removal Evaluation Report proposed the following removal actions in response to the identified conditions:

- **Plugging the connections to Manhole A.** This action was intended to minimize the risk of hydraulic surcharge to the pipe plug, thereby minimizing the risk of hydrocarbon releases from the pipe.
- Remove the accessible solid hydrocarbon material and place a cap over sediments
 containing hydrocarbon sheen in SG-04/SG-05 area. These actions were intended to
 minimize the risk of additional releases of hydrocarbons from this area to surface
 waters of the Port Washington Narrows and to prevent direct contact with these
 materials by beach users.
- **Install signage.** The purpose of the signs is to warn beach users about the presence of hydrocarbon contaminants in the beach sediments and provide agency contact information regarding the Site and the ongoing RI/FS process.

Upon completion of the removal evaluation, Cascade prepared a work plan describing the proposed removal actions in more detail. EPA approved the Final Removal Action Work Plan (Anchor QEA and Aspect, 2013b) and directed Cascade to perform the proposed removal actions (EPA, 2013c). After EPA's approval, Cascade implemented the removal action (2013 TCRA), which met all of the objectives specified in the Final Removal Action Work Plan including the following:

- Removing solid hydrocarbon material identified in the western beach area;
- Installing an organoclay mat and cover over the hydrocarbon sheen in subsurface sediments in the western beach area;
- Plugging Manhole A and the sump drain from the tank containment area;
- Completing monitoring inspections to confirm the effectiveness of the 2013 TCRA; and
- Installing required signage.

The work was completed in general accordance with the Final Removal Action Work Plan and documented in the TCRA Removal Action Report (Anchor QEA and Aspect, 2014). Three modifications to the scope of work specified in the Final Removal Action Work Plan were made with EPA approval based on the observed conditions:

- The organoclay mat and cover in the northeastern portion of the designed mat and cover area was extended to cover sediments exposed by the removal of the solid hydrocarbon material from the intertidal area.
- Manhole A was plugged by means of a concrete ring extending above the ground surface and capped with a bolted steel cover.
- Consistent with approvals from the City and pursuant to an access agreement with Penn Plaza Storage, LLC, a catch basin draining into the tank containment area was

rerouted to a City storm drain line to prevent accumulation of stormwater in the containment area.

Inspections of the 2013 TCRA areas were completed as specified in the Final Removal Action Work Plan (Anchor QEA and Aspect, 2013b). The inspections consisted of visual observation of the sediment cap and surrounding intertidal areas for the presence of product or sheen on the sediment or nearshore surface water and evaluation of the sediment cap for erosion. The inspections also included inspection of the manhole and tank containment areas to ensure that the plugs are intact and that surface water is not accumulating in these areas. Following completion of the removal action, inspections were performed at decreasing intervals over time, starting at weekly intervals and/or following significant precipitation events, and decreasing to quarterly intervals. Inspections have continued on a quarterly basis. To date, the constructed elements of the 2013 TCRA have been performing as designed with no surficial sheens observed in the organoclay mat area and no surface water accumulation in the manhole or tank containment areas. The constructed elements of the 2013 TCRA are shown in Figure 3-2.

3.4 Other Upland Investigations and Remedial Actions

Investigations and remedial actions conducted at other locations in the immediate vicinity of the Site may be relevant to characterizing the Site or understanding area-wide conditions. The only known upland investigations or remedial action performed in the immediate vicinity of the Site are those conducted at the Former SC Fuels Property.

Between 1997 and 2007, various consultants performed soil and groundwater sampling at the Former SC Fuels Property (Pacific Environmental, 1997; Noll, 1999 and 2000; GeoEngineers, 2002 and 2003; and GeoScience Management, 2007), including the following:

- Advancing 13 hand-auger borings, 18 direct-push soil borings, and 15 hollow-stemauger borings to a maximum depth of 22 feet;
- Installing 15 monitoring wells to a maximum depth of 20 feet;
- Collecting 12 soil confirmation samples during removal of four USTs; and
- Analyzing soil and groundwater samples for TPH, BTEX, and/or lead.

The investigations indicated the presence of TPH and BTEX in soil and groundwater on the Former SC Fuels Property and in the eastern portion of the Pennsylvania Avenue right-of-way. The TPH and BTEX concentrations exceeded Washington State Model Toxics Control Act (MTCA) Method A cleanup levels.

3.5 Other Sediment Investigations and Remedial Actions

In addition to the sediment data developed as part of previous investigations and remedial actions at the Site, other data sets have been compiled. The studies completed within the Port Washington Narrows and Dyes Inlet may provide information relevant to the RI/FS. Studies identified to date for these areas include the following:

• Chemical testing of sediments:

- 2008 and 2009 Puget Sound Assessment and Monitoring Program (PSAMP) Spatial/Temporal Monitoring, Central Sound (PSAMP, 2005 and 2009);
- 1989 to 2013 PSAMP Long-Term/Temporal Monitoring (PSAMP, 2005 and 2011a);
- 2009 PSAMP Urban Waters Initiative, Bainbridge Basin (PSAMP, 2005, 2009, and 2011b); and
- Ocean Survey Vessel *Bold* Summer 2008 Survey (USACE et al., 2009).
- Chemical testing of fish or shellfish tissue:
 - 2010 and 2012 Environmental Investment Project (ENVVEST) (Johnston et al., 2010; Brandenberger et al., 2012);
 - 2005 and 2007 NOAA Mussel Watch at station SIWP (Lauenstein and Cantillo, 1993; Kimbrough and Lauenstein, 2006; Kimbrough et al., 2006; and Kimbrough et al., 2008); and
 - o 2001 303d Ecology clam and crab sampling data (Ecology, 2002).
- Studies of surface water quality:
 - An Integrated Watershed and Receiving Water Model for Fecal Coliform Fate and Transport in Sinclair and Dyes Inlets, Puget Sound, Washington (Johnston et al., 2009); and
 - Sinclair and Dyes Inlets Fecal Coliform Total Maximum Daily Load:
 TMDL and Water Quality Implementation Plan (Lawrence et al., 2012).
- Regional studies of contaminant source inputs to these water bodies:
 - Contaminant Mass Balance for Sinclair and Dyes Inlets, Puget Sound, Washington (Crecelius et al., 2003).

Evaluation of this sediment and tissue data is discussed further in Section 3.9.

3.6 Existing Data and Data Usability

The existing Site characterization data have been reviewed in terms of data usability for the RI/FS. The existing data include data for the Former Gas Works Property and also data for sediments and tissue within the Port Washington Narrows, Dyes Inlet, and nearby portions of Puget Sound.

Data quality review included the definition of minimum data acceptability criteria (MDAC). Relevant guidance was applied, including the following:

- EPA (1988a) Guidance for Conducting Remedial Investigations and Feasibility Studies Under CERCLA;
- EPA (1992) Guidance for Data Useability in Risk Assessment, Part A;
- EPA Contract Laboratory Program Functional Guidelines for Data Review (variable dates for different analyte groups); and
- EPA (2009) Guidance for Labeling Externally Validated Laboratory Analytical Data for Superfund Use.

3.6.1 Minimum Data Acceptability Criteria

The MDAC evaluations of historical soil, groundwater, and sediment investigations are described for each sampling event in Table 3-8. ¹² MDAC evaluations of existing sediment and tissue investigations are described in Table 3-9. This MDAC review considered the following criteria:

- Work Plan Documentation:
 - O Documentation describing the sampling program or event, the methods used, and the parties involved in sample collection must be available.
 - Collection methods must be clearly defined and be adequate for obtaining representative and quantitative information.
 - The purpose of data collection should be available.
- Sample Location and Collection Methods:
 - Sample coordinates and a qualitative understanding of accuracy (i.e., knowledge of how the location was established or the method by which the coordinates were obtained) must be documented. The coordinate system must be documented.
 - Sample collection method and matrix must be documented. For example, a water sample must be identified as to whether it is a surface water, porewater, or groundwater sample and whether it is whole water or filtered (i.e., total versus dissolved fraction). Temporal or spatial compositing and sample volume must be identified. For tissue samples, tissue preparation must be documented.
 - Sample depths and, where applicable, start and end depths must be identified.
 - Sample storage methods must be documented and consistent with approved methods, including holding time and preservation.
 - Sample chain of custody must be documented.
- Laboratory Analysis:

O Data tables are available (not in summary format) with laboratory reports and data validation information

- Appropriate detection limits and quantitation limits are achieved so that the data meet the RI data quality objectives (DQOs) for environmental investigations:
 - Detection limits, units for each detection limit, and data qualifiers must be reported. Nondetected results must have the associated detection or reporting limits indicated. Data qualifiers must follow EPA guidance or be defined in documentation.
 - Analytical methods must be documented and acceptable based on EPA guidance.

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¹² Investigations conducted under the Order for the Site and performed in accordance with EPA-approved Quality Assurance Project Plans (i.e., the 2013 TCRA) are not included in the MDAC tables.

- Measurement instruments and calibration procedures must be documented.
- Toxicity and bioaccumulation test methods must be documented, including any deviations from standard protocols. For risk assessment, test methods must follow standard protocols, including controls and reference tests. Proper documentation to assess methods and statistical treatment must be available. Where possible, statistical results should be recalculated from the raw test data.
- Taxonomic data must be reported to the lowest practicable taxonomic level on a sample-specific basis, with scientific nomenclature. Taxonomic levels must be sufficient to assess relevant metrics for ecological risk assessment (ERA), such as feeding guilds or stress-induced compositional changes in the community.
- Collection methods, sample preservation, and sample preparation methods must be documented.
- Biological community metric calculations must be defined and documented.
- Quality Control and Data Validation:
 - Documentation of field and laboratory quality control samples (duplicates and blanks) must be present.
 - o Analytical chemical data must have been validated and qualified consistent with EPA functional guidelines or EPA Region 10 validation practices.
 - Hard copies of laboratory data reports (e.g., Form 1 or Certificates of Analysis) must be available to verify that electronic or tabulated data were accurately transcribed or transmitted.

3.6.2 Data Usability

Based on the results of the MDAC evaluation and considering the data representativeness for current Site conditions, the data were classified in one of the following data usability (DU) categories:

- **DU-1.** These data meet most or all of the MDAC requirements and are considered reasonably representative of Site conditions. DU-1 data are used in this Work Plan for COPC and source identification and preliminary evaluations of the nature and extent of contamination. These data may also be useful in the identification of data gaps and data needs, such as the mussel, clam, and crab results presented in this Work Plan.
- **DU-2.** These data meet most of the MDAC requirements but have been superseded by more current or higher quality data for representation of the nature and extent of contamination. If no DU-1 data is available (e.g., 2010 surface sediment results underlying the TCRA cap.), then DU-2 data are used in this Work Plan for COPC identification, source identification, and the preliminary evaluations of the nature and extent of contamination.

• **DU-R.** These data do not meet the MDAC requirements and are not used in this Work Plan.

Of the existing data, the data were classified as follows:

- **DU-1**:
- o All data collected during the 2013 TCRA.
- Soil data, sediment data for analytes other than PAHs, and groundwater data from monitoring wells, collected during the 2008 TBA.
- Soil and groundwater data collected during the 2007 Preliminary Upland Assessment. These data met most of the MDAC criteria but underwent minimal data validation.
- Regional sediment monitoring data collected under the following programs:
 - 2008 and 2009 PSAMP Spatial/Temporal Monitoring, Central Sound;
 - 1989 to 2013 PSAMP Long-Term/Temporal Monitoring;
 - 2009 PSAMP Urban Waters Initiative, Bainbridge Basin; and
 - Ocean Survey Vessel Bold Summer 2008 Survey.
- o 2010 and 2012 ENVVEST mussel data.
- 2005 and 2007 NOAA Mussel Watch at station SIWP.
- 2001 303d Ecology clam and crab sampling data.
- DU-2:
- Sediment data collected during the 2010 TCRA and sediment data for PAHs collected during the 2008 TBA. These data met most of the MDAC criteria but have been superseded by DU-1 data collected in 2013, after the 2010 TCRA was completed.
- DU-R:
- Soil and sediment data collected during the 1995 Ecology Field Inspection. These data had limited documentation, including poorly documented sampling locations, no documentation of collection or sample handling methods, and no chain of custody.
- Groundwater data collected from temporary borings during the 2008 TBA. The samples were not filtered, and the data are not considered representative of groundwater conditions because of potential bias due to sample turbidity.

3.7 Existing Data Summary

This Section summarizes existing relevant data for soil, groundwater, and sediment. The data have been used to prepare the preliminary CSM (Section 4) to support the definition of the Initial Study Area (see Section 5.1) and to develop the scope of work for the RI. The existing data will be used in the RI to help assess the nature and extent of contamination. They include data from the 2007 Preliminary Upland Assessment, select data from the 2008 TBA, and data from the 2013 TCRA. Data classified as DU-1 (see Section 3.6) are

included in the tables and figures associated with this Section. Data summary tables for each medium that include all data classified as DU-1 or DU-2 are provided in Appendix F.

3.7.1 NAPL Occurrences

Based on historical operations, LNAPL and/or DNAPL may be present at the Site. Gasoline and diesel petroleum products (LNAPLs) were transferred from docks via pipelines and stored in tank farms on the Former Gas Works Property and on three adjacent bulk fuel storage properties. Light oil (LNAPL) was generated as a byproduct of manufactured gas production and was stored in the south central portion of the Former Gas Works Property. Tars (generally DNAPLs) generated as a byproduct of manufactured gas production were potentially generated at several locations along the gas production process (including the scrubbers, gas holder, and purifier) and were reportedly managed/stored/placed in several areas of the Site, including: the Former Ravine fill area; the tar wells and residue cistern, adjacent to the Former Ravine fill area; a drain pipe suspected to be the former outfall that was removed and plugged during the 2010 TCRA; and the former tar pit reportedly located in the southwestern corner of the Former Gas Works Property.

Previous field investigations have included observations of NAPL (e.g., product or oil droplets) or indicators of the potential presence of NAPL (e.g., heavy sheens or staining). NAPL has been observed at the Site in soil and sediments at several locations, as follows:

- In shallow intertidal sediment in the vicinity of the stormwater pipe outfall that was removed, plugged, and overlaid with an organoclay mat during the 2010 TCRA (see Section 3.3.1).
- In an area of shallow intertidal sediment north of the former gas works that was overlaid with an organoclay mat during the 2013 TCRA (see Section 3.3.2).
- At the following soil borings:
 - MW-3 at a depth of 5 feet ("dark staining, creosote-like/solvent odor").
 This location is on the Former Gas Works Property in the vicinity of former tanks reportedly used to store tar.
 - MW-4 at a depth of 30 feet ("strong gas/diesel odor, product observed on grains¹³"). This location is on the Sesko Property downgradient of the former petroleum tank containment area.
 - MW-6 at a depth of 2 feet ("creosote-like odor and black tar-like substance) and 15 feet ("sheen with dark black creosote-like staining").
 This location is within the footprint of the former gas holder.
 - SP-03 at depths of 5 feet and 8 feet ("black coated sand...oil materials"),
 and 13.5 feet ("black coated sand...saturated with oil material"). This

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¹³ The exact same description of product noted at MW-4 was also noted at the same depth (30 feet) on the log for MW-6, but was in conflict with other field observations at the 30-foot depth interval at MW-6 (no sheen/slight sheen and low PID). Therefore, it is assumed that the note on MW-6 at 30 feet is an error.

location is in the Former Ravine fill area adjacent to the former residue cistern and tar wells.

No NAPL samples have been collected or submitted for testing to characterize its chemical or physical properties. Sediments containing heavy sheens that were sampled during the 2010 and 2013 TCRAs exhibited higher concentrations of PAH compounds than surrounding sediment samples.

Other potential indicators of NAPL presence include very high concentrations of organic compounds in soil (i.e., close to or above potential residual saturation levels) or in groundwater (e.g., greater than 10 percent of a component's aqueous solubility). Naphthalene has been detected in one well (MW-4, located in the Former Ravine fill area) at greater than 10 percent of its solubility.

3.7.2 Soil Data

Soil samples were collected as part of the investigations conducted in 2007, 2008, and 2013. These data sets include the following:

- **2007.** Seventeen soil samples were collected from eight explorations and analyzed for TPH, metals, SVOCs (including PAHs), VOCs and PCBs.
- **2008.** Forty-three soil samples were collected from eight explorations and analyzed for TPH, metals, SVOCs (including PAHs) and VOCs.
- 2013. Two soil samples were collected from two explorations and analyzed for PAHs.

Table 3-10 summarizes the number of samples collected for analysis of each constituent and an evaluation of detected concentrations as compared to the initial PRG. Data for metals are also compared to the regional natural background concentrations established by Ecology (Ecology, 1994). The soil analytical data are summarized in tables provided in Appendix F.

The constituents detected in soil at concentrations above the initial PRGs include the following:

- VOCs, including 1,2,4-trimethylbenzne, benzene, and ethylbenzene;
- 2-methylnaphthalene;
- PAHs; and
- Metals, including antimony, arsenic, cadmium, chromium, cobalt, copper, lead, manganese, mercury, nickel, thallium, vanadium, and zinc.

Other than PAHs, no SVOCs were detected at concentrations above the initial PRGs except for a single detection of 2-methylnaphthalene; however, the reporting limits for a subset of SVOCs exceed the initial PRGs at some locations (Table 3-10 and Appendix F). Practical quantitation limits (PQLs) for COPCs based on standard analytical methods are provided in Table 3-10 for comparison.

PCBs were not detected in soil; the reporting limits for PCBs in all samples were less than the initial PRGs (Appendix F).

Initial PRGs are not identified for TPH, which is not a hazardous substance under CERCLA. However, identifying the nature and extent of different TPH products (e.g., gasoline or diesel) may be helpful in defining contaminant sources. TPH data should be used with caution at sites, such as MGP sites, where non-petroleum hydrocarbon mixtures are present (e.g., carbureted water-gas tar). Therefore, an understanding of the type of product present, as assessed by sample chromatogram review or forensic analysis and interpretation, is needed to correctly interpret TPH data. For the purposes of this Work Plan, TPH distribution was not evaluated but will be evaluated in the RI.

A summary of VOCs, PAHs, and metals detected at concentrations above the initial PRGs is provided in the following Sections by analyte group. The maximum concentration detected at each boring location and a comparison to the initial PRGs and/or natural background concentrations in shallow soil (0 to 10 feet deep) and deeper soil (greater than 10 feet deep) is provided for the primary constituents detected at concentrations greater than the initial PRGs¹⁴ (Figures 3-3 through 3-14). As described in Section 3.1.2, initial PRGs for surface soil include a consideration of potential terrestrial ecological exposure, whereas the initial PRGs for subsurface soil do not. For the purposes of presenting existing data in this Work Plan, soil data in the 0 to 10 feet in depth is compared to initial PRGs for surface soil to account for potential terrestrial ecological exposures. ¹⁵

3.7.2.1 Volatile Organic Compounds

Two BTEX compounds, benzene and ethylbenzene, were detected at concentrations greater than the initial PRGs. The most frequent detections of benzene above the initial PRG occurred at two locations: in shallow soil collected at sample location MW-3, in the vicinity of the former finished gas storage tanks, and at sample location SP03, near the edge of the Former Ravine (Figure 3-3). Benzene was not detected in any deeper soil samples at a concentration above the initial PRG (Figure 3-4). BTEX compounds are potentially an indicator of MGP-related releases but may result from other sources (e.g., gasoline-range TPH or industrial solvents).

Concentrations of 1,2,4-trimethylbenzene were detected above the initial PRG in four soil samples. 1,2,4-trimethylbenzene is a component of carbureted water-gas tar and petroleum.

3.7.2.2 Polycyclic Aromatic Hydrocarbons

The maximum concentrations of naphthalene in shallow and deeper soil are shown on Figures 3-5 and 3-6, respectively. The concentrations of total carcinogenic PAHs (cPAHs)¹⁶ in shallow and deeper soil are shown on Figures 3-7 and 3-8, respectively. The

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¹⁴ Primary constituents shown on the figures include those detected in excess of the PRGs and the natural background concentrations with the greatest frequency or magnitude.

¹⁵ As noted in Section 4.3, 'surface soil' for the purposes of the ecological and human health risk assessments may vary depending on the applicable exposure pathway but is generally 3 feet or less in depth. However, there is limited existing data in this interval; therefore, separate figures for surface soil were not prepared for this Work Plan.

¹⁶ Concentrations of total cPAHs are provided in benzo(a)pyrene toxic equivalent concentrations (EPA, 1993).

vertical distribution of naphthalene concentrations in soil is illustrated along geologic cross sections A–A′, B–B′, C–C′, and D–D′ on Figures 2-9 through 2-12, respectively.

The concentrations of total cPAHs and naphthalene exceeding the initial PRGs were detected at sampling locations that correspond to operational areas of the former gas works. In shallow soil, the highest concentrations of both total cPAHs and naphthalene were detected at sample location MW-3, advanced in the vicinity of the storage tanks, which held light oil and carbureted water-gas tar (Simonson, 1997b). Likewise, the highest concentrations of both total cPAHs and naphthalene in deeper soil were detected at sample location MW-6, which was advanced at the location of the former gas holder.

Generally, concentrations of naphthalene and cPAHs on the Former Gas Works Property are highest in shallow soil and decrease with depth (MW-3 and SP03, for example). However, at MW-6, advanced at the location of the former gas holder, PAH concentrations detected in deeper soil were much higher than those in shallow soil. Because the gas holder was reportedly at least 10 feet deep, this finding may indicate that the gas holder was filled with cleaner soil after it was demolished. Also, the concentrations of PAHs detected in deeper soil were greater than those in shallow soil at well MW-8, located hydraulically downgradient of the former gas works operational area.

The concentrations of total cPAHs exceeding the initial PRG have been detected in soil samples collected between depths of 3 and 40 feet. The highest concentrations of total cPAHs were detected in shallow soil, between the depths of 5 and 12 feet, at well MW-3, well MW-6, and boring SP03 and in deeper soil at a depth of 25 feet at well MW-8.

The presence of cPAHs and naphthalenes is a potential indicator of MGP-related releases ¹⁷

3.7.2.3 Metals

The detectable concentrations or analytical reporting limits for a number of metals exceeded the initial PRGs. However, the concentrations of many of these metals did not exceed the natural background concentrations¹⁸ (Ecology, 1994):

- For manganese and antimony, all of the detected concentrations, and most of the reporting limits, are below the background concentrations. ¹⁹
- Cobalt and vanadium were detected in all of the soil samples analyzed for metals, with many concentrations exceeding the initial PRGs; however, the detected concentrations are generally within the range of regional background concentrations.
- Thallium was detected at concentrations above the initial PRGs in most of the soil samples analyzed; a natural background concentration for thallium was not available.

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¹⁷ Carcinogenic PAHs and naphthalenes can also originate from other sources, including petroleum hydrocarbons or creosote. Forensic analyses, such as PAH fingerprinting, may be useful during the RI to help distinguish and identify potential sources of contamination.

¹⁸ Puget Sound background concentrations of metals were used for screening when available. When not available, Washington State background concentrations were used.

¹⁹ The Puget Sound regional background concentration for antimony has not been researched. The referenced background concentration is based on regional data from the Spokane Basin.

Detected concentrations of cadmium, lead, and zinc are within the range of regional background concentrations at most sample locations, except for borings MW-5, MW-8, and SP03, which are located at the northeast corner of the Former Gas Works Property in the shoreline and Former Ravine fill area.

Arsenic, chromium, copper, and nickel were detected at concentrations above the initial PRGs and background concentrations at several locations. Figures 3-9 through 3-14 depict the concentrations of arsenic, copper, and nickel²⁰ in shallow and deeper soil. Concentrations of these metals in deeper soil do not exceed the initial PRGs, with the exception of arsenic, which was detected at a concentration above the initial PRG but below the natural background concentration. Concentrations of arsenic, copper, and nickel in shallow soil exceed the initial PRGs and the natural background concentrations at several locations. Arsenic was detected at concentrations above the natural background concentration at two locations: SP03 (Former Ravine fill area) and MW-3 (within the footprint of former gas works operations and the current industrial park). Copper, chromium, and nickel were sporadically detected across the Former Gas Works Property at concentrations above the natural background concentrations, but their maximum concentrations were only slightly above their respective background concentrations (62.7 milligrams per kilogram [mg/kg] versus 38 mg/kg for copper; 60.8 mg/kg versus 48 mg/kg for chromium; and 60.9 mg/kg versus 48 mg/kg for nickel). The sources of these exceedances are unclear from the existing data. Possible sources include contaminated fill. historical industrial operations, or natural background variability.

3.7.3 Groundwater Data

Groundwater samples were collected as part of the investigations conducted in 2007 and 2008. Groundwater samples were collected and analyzed for petroleum hydrocarbons, metals, SVOCs including PAHs, VOCs, and PCBs. Table 3-11 summarizes the number of samples collected for analysis of each constituent and the results of a comparison of detected concentrations to the initial PRGs, which include concentrations protective of groundwater and surface water. The groundwater analytical data are provided in Appendix F.

The constituents detected in groundwater at concentrations greater than the initial PRGs include the following:

- Metals: arsenic, beryllium, chromium (both total and hexavalent), cobalt, copper, lead, manganese, mercury, nickel, thallium, vanadium, and zinc;
- PAHs: acenaphthene, benzo(g,h,i)perylene, dibenzofuran, fluoranthene, florene, phenanthrene, pyrene, naphthalenes, and total cPAHs;
- Pentachlorophenol (PCP); and
- VOCs: benzene, ethylbenzene, xylenes, 1,2,4- and 1,3,5-trimethylbenzene, 1,2-dichloroethane, carbon tetrachloride, chloroform, isopropylbenzene, n-hexane, and trichloroethene.

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²⁰ Arsenic, copper, and nickel were mapped in soil because these constituents were also most frequently detected in groundwater at concentrations greater than the surface water or groundwater initial PRGs.

Other than the above-listed constituents, no SVOCs or VOCs were detected at concentrations above the initial PRGs; however, the reporting limits for a subset of SVOCs and VOCs exceed the initial PRGs at a number of locations (Table 3-11 and Appendix F). PCBs were not detected in groundwater; however, the reporting limits for PCBs in all samples were above the groundwater initial PRG (Appendix F).

The existing groundwater data are limited, with one sampling event at 10 locations and no groundwater data collected since 2008. The data are useful for the preliminary identification of COPCs, and they indicate where groundwater impacts may be located. Some of the existing data were collected from wells that are still in place. These wells can likely be used for future monitoring, and the comprehensive data set will likely be useful in evaluating long-term trends in groundwater quality.

VOCs, PAHs, PCP, and metals detected at concentrations above the initial PRGs are discussed in the following Sections by analyte group. The concentration detected at each monitoring well and a comparison to the groundwater initial PRGs are provided for the primary constituents detected at concentrations above the initial PRGs²¹ on Figures 3-15 through 3-19.

3.7.3.1 Volatile Organic Compounds

One or more of the BTEX compounds were detected in groundwater samples collected at all of the monitoring wells, except for wells MW-1 and SP02. The detected concentrations of benzene in groundwater are shown on Figure 3-15. The highest concentrations were detected in wells MW-3, MW-6, and MW-8 (in and downgradient of the former gas works operation area).

3.7.3.2 Polycyclic Aromatic Hydrocarbons

Detected concentrations of total cPAHs were above the initial PRGs in groundwater samples collected from wells MW-3 through MW-8 (Figure 3-16) located on the Former Gas Works Property. The highest concentration of total cPAHs in groundwater was detected at well MW-4. There were no detected concentrations of cPAHs in the groundwater samples collected from wells MP04, SP02, MW-1, and MW-2.

The results for other PAHs are the following:

- Dibenzofuran and pyrene were detected at concentrations above the initial PRGs in the groundwater sample collected from well MW-4; and
- Naphthalenes, including 1-methylnaphthalene and naphthalene, were detected in groundwater samples collected from wells SP02, MP04, MW-3, MW-4, MW-5, MW-6, MW-7, and MW-8 at concentrations exceeding the initial PRGs. The highest concentrations of naphthalene were detected at wells MW-4 and MW-8 (Figure 3-17).

3.7.3.3 Pentachlorophenol

PCP was detected in groundwater at a concentration exceeding the groundwater and surface water initial PRGs at well MW-8.

²¹ Primary constituents shown on the figures include those detected with the greatest frequency or magnitude above the groundwater initial PRG.

3.7.3.4 Metals

The highest concentrations of metals in groundwater were generally detected at wells MW-3 and MW-4. MW-3 is located in the central portion of the Former Gas Works Property, in the vicinity of the former finished gas storage tanks and former metal finishing operations. MW-4 is located within the Former Ravine fill area, in the central portion of the Sesko Property. Results for specific metals are the following:

- Arsenic was detected in all of the groundwater samples analyzed, at concentrations ranging from 0.6 to 26 micrograms per liter (μg/L), all of which exceed both the groundwater initial PRG and the surface water initial PRG. Figure 3-18 depicts the concentrations of arsenic in groundwater, which are highest in the central portion of the Former Gas Works Property, at wells MW-3 and MW-4
- Hexavalent chromium was detected in groundwater samples collected from wells MW-1 and MW-3 through MW-8 at concentrations exceeding the groundwater initial PRG. The concentrations detected in wells MW-5 and MW-8 also exceed the surface water initial PRG. Figure 3-19 depicts the concentrations of hexavalent chromium in groundwater.
- Total chromium and lead were detected in groundwater at concentrations above both the groundwater initial PRGs and the surface water initial PRGs in the samples collected from wells MW-3 and MW-4.
- Copper and nickel were detected at concentrations exceeding the surface water initial PRGs at most of the sampling locations; none of the concentrations of copper and nickel exceeds the groundwater initial PRGs. The highest concentrations of copper and nickel were detected in groundwater samples collected from wells MW-3 and MW-4.
- Concentrations of cobalt, manganese, thallium, and vanadium exceeding the groundwater initial PRGs were detected in the groundwater sample collected from well MP04.

3.7.4 Sediment Data

Available sediment data for the Site include those collected in 2008 as part of the TBA, in 2010 as part of the 2010 TCRA, and in 2013 as part of the 2013 TCRA. These data sets include the following:

- **2008.** Five surface sediment samples from the beach north of the Former Gas Works Property were analyzed for TPH, VOCs, SVOCs, and metals.
- **2010.** Thirty-two surface sediment samples collected during the 2010 TCRA were analyzed for VOC and SVOCs.
- 2013. Thirty-nine surface sediment samples collected during the intertidal sediment sampling program were analyzed for total solids (TS), total organic carbon (TOC), and SVOCs.
- **2013.** Seventeen subsurface sediment samples were collected by direct-push methodology at seven locations (boring depths ranged from 4- to 5-feet below the sediment surface). Samples from 4 discrete intervals were analyzed for VOCs, and samples from 17 subsurface intervals were analyzed for TS, TOC, and SVOCs.

Table 3-12 presents these sediment data and concentrations relative to the initial PRGs identified in Section 3.1.2. Where applicable, reference values are also presented for natural background concentrations of contaminants in Puget Sound sediments or soils.

Figures 3-20 through 3-24 present the measured concentrations of PAHs in beach sediments at the Site. Data are presented on a dry-weight basis for benzo(a)pyrene, total low-molecular-weight PAHs (LPAHs), total high-molecular-weight PAHs (HPAHs), total cPAHs, and total cPAH toxic equivalent (TEQ) concentrations. The highest PAH concentrations were detected within and near the two removal action areas. East and west of these two areas, concentrations decrease rapidly.

3.8 Existing Data from Other Cleanup Sites

Soil and groundwater data collected on the Former SC Fuels Property include TPH, BTEX, and lead (Section 3.4). The majority of the soil data were collected prior to and during remedial actions (removal of USTs and surrounding contaminated soil), which occurred in 2002. The most recent groundwater monitoring data are from January 2007. During that sampling event, concentrations of benzene were detected in groundwater at concentrations up to 88 μ g/L on the Former SC Fuels Property and up to 49 μ g/L in the eastern portion of the Pennsylvania Avenue right-of-way (GeoScience Management, 2007). The extent of benzene detected in groundwater (detection limit 1 μ g/L) in 2007 is shown on Figure 3-25.

3.9 Data for Port Washington Narrows, Sinclair Inlet and Dyes Inlet

A number of high-quality sediment and tissue studies were identified for the Port Washington Narrows, Sinclair Inlet, and Dyes Inlet. The locations where sediment and tissue data with measured PAH concentrations were collected are shown on Figure 3-26. These data will not be used for data screening or COPC identification (see Section 4.4), but may provide useful information about conditions in the vicinity of the Site.

Because the delineation between the Site's contamination and other sources of contamination is not straightforward, these data were thought to be potentially useful to assess whether off-Site sediment quality could potentially affect conditions at the Site through sediment transport and recontamination processes. After review of these data, it was determined that additional sampling and assessment would be required to understand sediment transport into and out of the Site for the purposes of the FS. The approach being used to assess sediment transport mechanisms in the vicinity of the Site is described in Section 5.5.2.

No recently-collected water quality data for chemical contaminants within the Port Washington Narrows have been identified. Several studies have been conducted to assess potential contaminant inputs to Dyes Inlet and adjacent waters (Crecelius et al., 2003). The results of these and other available studies may be used qualitatively for evaluating the potential influence of nonpoint sources of pollution on the Site, but will not be relied upon for the baseline risk assessment.

4 Preliminary Conceptual Site Model

This Section presents the preliminary CSM, which has been developed based on available historical information, the current understanding of the environmental setting, and the findings of previous investigations, as presented in Sections 2 and 3. The CSM is a description of environmental conditions that includes sources of contamination, contaminant fate and transport in Site media, and potential routes of contaminant exposure for human and environmental receptors. A three-dimensional graphical CSM illustrating representative potential historical sources and migration of contaminants at the Site is provided on Figure 4-1, and a conceptual CSM cross section is shown on Figure 4-2. The CSM will be developed further during the RI and risk assessment as more Site-related information and data are gathered.

4.1 Potential Sources of Contamination

This Section summarizes potential sources of contamination on the Former Gas Works Property and on surrounding properties. The potential sources and locations associated with known and documented operations (both MGP and other) are presented in the following Sections; however, this discussion does not include undocumented or currently unknown potential sources or source areas, which may be identified through the collection and evaluation of data during the RI.

4.1.1 Former Gas Works Property Sources

Potential sources of contamination on the Former Gas Works Property include historical activities associated with the former gas works, as well as other activities on the property that are unrelated to gas works operations.

4.1.1.1 Sources Related to Gas Works Operations

The potential primary sources associated with the production of manufactured gas are depicted on Figure 2-3. The area in which the gas production process occurred is divided into potential source areas based on the predominant use and subsequent primary potential release mechanisms associated with each area. The primary potential source areas include the following:

- Coal/Coke Briquettes Area. As described in Section 2, solid feedstocks (coal and coke briquettes) were transported to the Former Gas Works Property by barge and offloaded and transported over the water, beach, and bluff to a concrete surface storage area in the northwest corner of the Former Gas Works Property. Coke briquettes have been observed on the beach and bluff, suggesting spills during the transport process. Additionally, coal/coke dust may have been swept off the concrete storage slab onto the surrounding ground surface.
- Tar and Petroleum Transfer Area. Petroleum products were delivered to the Former Gas Works Property and tar was removed from the Former Gas Works Property by barge. Petroleum and tar from pipelines along the dock and at the connection to the barges may have been released directly to sediment or surface water. A pipeline presumably ran between the dock and the byproduct storage area to transport tar to the dock, but the location is unknown.

- **Petroleum Storage Area.** Petroleum products were stored in ASTs in the northeastern portion of the Former Gas Works Property. The products reported to have been stored in these tanks include gasoline and diesel fuel oil. Transfer piping presumably ran from the storage tanks to the furnaces, but the exact location of transfer piping is unknown. Petroleum may have been released from tanks and piping to soil at the ground surface or shallow soil in this area.
- Gas Generation and Purification Area. The main process area was located in the central portion of the Former Gas Works Property and included the furnaces, scrubber, gas holder, and purifier. The primary potential sources associated with the gas works process consist of spills, drips, and leaks of spent liquids, oils, gas liquor, tar, and tarwater mixtures from aboveground equipment, piping, and storage tanks to the ground surface.
- Residuals Management Area. A map of the former plant shows tar wells and a residue cistern to the east of the purifiers. These were likely used for separation of tarwater emulsions prior to resale of the tar. The details of the tar wells and residue cistern are unknown, but they likely extended into shallow subsurface soil and may have either been lined or unlined at the base. A second area south of the main plant building was reportedly used for storage and/or separation of tar and tar-water emulsions in a tar pit. Oils and tar may have been released to the ground surface around these features or the shallow soil beneath them.
- Tar and Light Oil Storage Area. The southern portion of the Former Gas Works Property was used for the storage of tar and light oil in ASTs. Tar and light oil may have leaked or been spilled onto the ground surface in the vicinity of the ASTs. Finished gas may have contained small amounts of oil that condensed in the distribution piping and were collected in the drip tank. Light oil may have been released to shallow soil in the vicinity of the pipes and tank.
- Former Drainage Line Area. During the 2010 TCRA, a former drainage line on the Sesko Property that discharged to the Port Washington Narrows was identified. Tarlike hydrocarbons were identified in this drainage line, which was plugged during the 2010 TCRA (see Section 3.3.1). The alignment of the drainage line is similar to the alignment of a former City CSO outfall documented in historical records. Wastewater and associated contaminants may have discharged from this drainage line during and after operation of the former gas works.
- Ravine and Shoreline Fill Areas. Historical documents indicate gas works byproducts may have been placed into the western portion of the Former Ravine, to the east of the gas generation and purification area, and along the bluff to the north of the gas generation and purification area for some period of time. Materials that were reportedly placed along the shoreline include ash, cinders, slag, and soot. Materials that were reportedly placed in the Former Ravine include ash, cinders, slag, soot, spent scrubber media (tar-laden wood chips and shavings), and spent purifier filter media (wood chips and/or iron oxide). The approximate areas where gas works byproducts may have been placed are shown on Figure 2-3.

4.1.1.2 Sources Related to Other Operations on Former Gas Work Property

Other potential primary sources are associated with activities conducted after the shutdown and demolition of the former gas works, or they were conducted in the immediate vicinity of the former gas works. These sources are shown on Figure 2-4 and summarized as follows:

- **Bulk Petroleum Storage.** Petroleum products were delivered to Lent's at a dock offshore of the Sesko Property and stored in ASTs for distribution by fuel delivery vehicles. Petroleum may have been released from piping and storage tanks to the ground surface and/or shallow soil.
- Varied Light Industrial Use. Since the shutdown of the former gas works, the
 McConkey Property has been used for miscellaneous light industrial activities,
 including vehicle parking, metals fabrication, and equipment storage. Ecology
 inspections in 1992, 1993, and 1994 indicated poor housekeeping practices associated
 with some of these operations. These operations are potential sources of solvents,
 metals, and petroleum hydrocarbons, which may have been released to the ground
 surface as either solids (sandblast grit, paint sludges, etc.) or components of liquids.
- Equipment Storage and Repair and Debris Filling. In addition to the bulk
 petroleum storage described above, activities on the Sesko Property since the
 shutdown of the former gas works included boat maintenance and storage, automobile
 salvage, and equipment and debris storage. These activities may be sources of
 contaminants to soil, sediment, and surface water by direct discharge, dumping, or
 spills to the ground surface.
- Other Operations. Other operations have reportedly included filling of the Former Ravine and shoreline areas, particularly on the Sesko Property. These operations may have included disposal of incinerator refuse, garbage, and ashes; placement of concrete and piping debris; and/or placement of miscellaneous metal, concrete, and fiberglass debris associated with maintenance and salvage of boats and equipment. Fill placed along the shoreline and in the Former Ravine may have included materials that contained hazardous substances. Although the presence of fill material alone does not necessarily represent a contaminant source, hazardous substances associated with the fill may subsequently migrate to surrounding soil or groundwater.

4.1.1.3 Stormwater Discharge

Stormwater discharging to the Port Washington Narrows may contain contaminants and is a potential source of contamination to sediments or surface water. The outfalls that historically have captured or currently capture water at the Former Gas Works Property are the following:

• **Historical City Stormwater/CSO Outfall.** As noted in Section 4.1.1.1 (list item "Former Drainage Line Area"), a historical drainage line and outfall were located within and offshore of the Sesko Property. A section of the drainage line on the beach was reportedly removed by the City during installation of a force main in the 1990s. The drainage line was plugged and partially removed as part of the 2010 TCRA (see Section 3.3.1). An upland manhole and storm drainage lines believed to be connected historically to the drainage line were plugged as part of the 2013 TCRA.

• **McConkey Drainage Line.** A small drainage line discharges stormwater from a shallow catch basin on the McConkey Property to the Port Washington Narrows.

4.1.2 Sources Related to Operations on Adjacent Properties

Potential primary sources on adjacent properties include the following:

- **Bulk Petroleum Storage.** Petroleum products were delivered by barge to bulk fuel storage facilities at the Former ARCO Dock, the Former Sesko Dock, and the former SC Fuels Dock and stored in ASTs or USTs for distribution by fuel delivery vehicles. These petroleum storage facilities were located on the Former ARCO Property located west of the former gas works and the Former SC Fuels Property. Petroleum may have been released from piping and storage tanks to the ground surface and/or shallow soil while these operations were ongoing.
- Varied Light Industrial Use. The Penn Plaza Property has been used for miscellaneous light industrial activities, including spray painting, a pipe shop, vehicle parking for a petroleum distributor, truck repair electroplating, metals fabrication, and equipment storage. Ecology inspections in 1992, 1993, and 1994 indicated poor housekeeping practices associated with some of these activities. These activities are potential sources of solvents, metals, and petroleum hydrocarbons, which may have been released to the ground surface as either solids (sandblast grit, paint sludges, etc.) or components of liquids.

4.1.2.1 Stormwater Discharge

A number of documented stormwater and CSO outfalls are located within the Port Washington Narrows and Dyes Inlet (Section 2.7), including the two outfalls described in Section 4.1.1.3. Other nearby outfalls or discharge lines include the following:

- Current City Stormwater/CSO Outfall. An active City stormwater/CSO outfall is located along the Port Washington Narrows, offshore of the end of Pennsylvania Avenue. This outfall is located adjacent to the 2010 TCRA area (Figure 3-2).
- **Drain Line.** A drain line from an oil-water separator on the Former SC Fuels Property discharges to the Port Washington Narrows.

4.2 Contaminant Migration and Transformation

Contaminants derived from the sources described in Section 4.1 may have been released to soil (surface and shallow subsurface), sediment, and/or surface water. Representative potential releases (e.g., leaks or spills from equipment, tanks, or piping; placement of contaminated fill materials; and discharges from outfalls) are shown conceptually on Figures 4-1 and 4-2. The released contaminants may have migrated from one location to another or from one medium to another. Contaminants may also undergo attenuation or transformation processes within media. The contaminant migration pathways and transformation processes are described in the following Sections.

4.2.1 Migration Pathways

Examples of potential contaminant migration pathways between media are shown conceptually on Figures 4-3, 4-4, and 4-5, and include the following:

- Migration of contaminants from surface soil to subsurface soil (e.g., leaching or product migration);
- Contaminant leaching or NAPL migration from soil/NAPL to groundwater;
- Groundwater/NAPL transport within the saturated zone;
- Groundwater discharges to surface water;
- Contaminant partitioning between groundwater and sediments (including sediment porewater);
- Migration of volatile NAPL/soil/groundwater contaminants to air;
- Migration of surface soil contaminants as fugitive dust;
- Release of surface soil contaminants to stormwater;
- Uptake of contaminants by terrestrial or aquatic biota; and
- Migration of contaminated sediments by sediment transport.

Based on the data collected to date (see Section 3.7), contaminants have been identified in soil, groundwater, and sediment. No Site-specific surface water, air, or tissue data are available. Contaminant occurrences in these media may be due to direct releases or subsequent migration, for instance:

- Soil contamination may be the result of contaminated fill materials, downward flows of NAPL releases²² through the subsurface and the coating of soil grains, or sorption of contaminants from other media (e.g., soil vapor, infiltrating stormwater, or groundwater).
- Groundwater contamination may be the result of direct discharge of contaminated
 aqueous materials and their migration downward through the subsurface and mixing
 with groundwater, leaching of NAPL in contact with groundwater, or stormwater
 infiltration of the subsurface, leaching of contaminants from NAPL or contaminated
 soil, and contaminant mixing with groundwater.
- Contaminants in sediment may be the result of direct releases to surface sediments
 (e.g., documented discharges from outfalls, undocumented spills, or leaks from dock
 piping and transfer operations); subsurface migration of contaminated groundwater or
 NAPL from the uplands, and migration through sediments; or a combination of
 sources. In particular, two sediment "hot-spot" areas were addressed by the 2010 and
 2013 TCRAs:
 - The 2010 TCRA addressed a drain pipe that contained residual NAPL and surrounding contaminated sediments, which appeared to be the primary source of contamination in this area. The historical and ongoing contribution to sediment contamination from other potential sources in this area, including groundwater discharge, stormwater runoff, and the City CSO, is unknown.

²² Liquid releases generally move downward, through the subsurface by means of gravity, but they may move laterally by preferential migration pathways if a barrier (e.g., low-permeability soils or, for NAPLs that are less dense than water, groundwater) is encountered.

The 2013 TCRA addressed an area of heavy sheen located in shallow subsurface sediments and solid surficial material containing high PAH concentrations. It is likely that the solid surficial material, which would be immobile in the subsurface, was placed at or near its locations; however, the source of the material is unknown. The source of the subsurface sheen is also unknown. During the TCRA investigation, a sheen was observed up to the base of the bluff. However, there are insufficient data to determine whether this contamination is contiguous with contamination in the upland.

Representative migration pathways, including subsurface migration pathways, are indicated on Figures 4-1 and 4-2.

4.2.2 Transformation Processes

In addition to contaminant migration pathways, contaminant concentrations in media can be reduced or attenuated by various combinations of natural processes. Examples of such processes include the following:

- Chemical or biological degradation of contaminants in soils, groundwater, sediments;
- Tidally induced mixing of groundwater near the groundwater/surface water interface;
- Natural recovery of marine sediments by burial, mixing, and/or degradation processes; and
- Metabolic transformation or elimination of chemical contaminants from the tissues of upland or aquatic biota.

4.3 Exposure Pathways and Receptors

Exposure pathways are the routes through which people or ecological organisms are exposed to chemicals (e.g., through eating, drinking, breathing, touching). Relevant and representative human and ecological receptors that may use the Site are summarized in Figures 4-3, 4-4, and 4-5. These figures illustrate how humans and ecological receptors may be exposed to chemicals. To determine whether an exposure pathway is complete and, therefore, exposure can occur, the following four elements must be evaluated:

- Source of chemical release;
- Release or transport mechanism (or media in cases involving media transfer);
- Exposure point (a point of potential human or ecological contact with the contaminated exposure medium); and
- Exposure route (e.g., ingestion or dermal contact) at the exposure point.

If any of these elements are missing, the pathway is considered incomplete and exposure will not occur. The definitions of all possible exposure pathway designations are as follows:

• Preliminary Complete Exposure Pathway – There is a source, a release and transport mechanism from a source, an exposure point where contact can occur, and an

- exposure route through which contact can occur. These complete exposure pathways will be quantitatively evaluated in the risk assessment.
- Preliminary Complete Exposure Pathway, Low Exposure Potential There is a source, a release and transport mechanism from a source, an exposure point where contact can occur, and a limited exposure route through which contact can occur. These complete exposure pathways will be quantitatively evaluated in the risk assessment.
- Currently Incomplete Exposure Pathway, Potential Future Exposure Evaluation –
 There is a source, a release and transport mechanism from a source, an exposure point
 where contact can occur, and a currently incomplete exposure route through which
 contact can occur. If future conditions change, the exposure would be complete. These
 exposure pathways will be quantitatively evaluated in the risk assessment.
- Preliminary Incomplete Exposure Pathway There is either no source, no release or transport mechanism from a source, no exposure point where contact can occur, and/or no exposure route through which contact can occur for the given receptor. Pathways considered incomplete will not be evaluated further in the risk assessment.

Figure 4-3 illustrates different exposure pathways that could affect people using the Site or nearby areas. The potential exposure of people to Site-related COCs differs in terms of both how those people use the Site and which areas of the Site are used. (i.e., beach/aquatic areas and upland areas). Some land uses could also change over time. For example, the Site is not zoned for residential land use, but as part of the risk assessment activities, it may be prudent to evaluate potential future residential land use to understand the implications of changes in land use or zoning. Similarly, shellfish harvesting in the Port Washington Narrows is restricted due to shellfish harvesting closures unassociated with the former gas works. However, it may be prudent to evaluate potential future shellfish harvesting to understand potential exposures should those shellfish harvesting restrictions be lifted.

Preliminary complete current and future human exposure pathways to contaminated media include dermal contact with and incidental ingestion of soil or sediment, dermal contact with groundwater, inhalation of fugitive dust and vapors, and consumption of fish/shellfish that are potentially contaminated with bioavailable Site-related contaminants. Preliminary incomplete current and future human exposure pathways will be evaluated further as part of the RI and risk assessment (see Section 6 for planned RI and risk assessment methodology). The preliminary human exposure scenarios relevant to the Site include the following:

- Human Use of Beach/Aquatic Site Areas:
 - Recreational Beach Users. There is a potential for limited recreational beach use by individuals residing near the Site. During recreational use of the beach, these individuals could be exposed through dermal contact and incidental ingestion of sediment and porewater.
 - Tribal Subsistence and Recreational Consumers of Fish/Crab from the Port Washington Narrows. The portions of the Port Washington Narrows adjacent to the Former Gas Works Property currently support the collection and consumption of fish and crabs under WDFW regulations. The Port Washington Narrows is also a Usual and Accustomed area of the

- Tribe. Consumers of fish and crabs may be exposed to Site-related COCs through direct contact with and incidental ingestion of sediment and porewater during harvesting activities, and through ingestion of fish/crab tissue.
- O Tribal Subsistence and Recreational Consumers of Shellfish at the Site (currently restricted by Shellfish Harvesting Closures). The portions of the Port Washington Narrows adjacent to the Former Gas Works Property are currently closed to shellfish harvesting by WDOH (due to water quality concerns associated with CSOs and other non-Site-related concerns); however, exposures associated with shellfish harvesting will be evaluated to understand potential future risks should the shellfish harvest restrictions be lifted. Future consumers of shellfish may be exposed to Site-related COCs through ingestion of shellfish tissue and dermal contact with and incidental ingestion of sediment and porewater during harvesting activities.
- Beach Construction/Excavation Workers. This scenario relates to workers
 performing utility upgrades or maintenance or other activities that involve
 the disturbance of sediment in the beach area adjacent to the Former Gas
 Works Property. Beach construction workers could be exposed to Siterelated COCs through direct contact with porewater and through direct
 contact with and incidental ingestion of surface and subsurface beach
 sediment
- Human Use of Upland Site Areas:
 - Occupational Workers. The Former Gas Works Property and the properties in the vicinity are zoned for industrial uses. Occupational workers at the Site could be exposed to Site-related COCs through direct contact with and incidental ingestion of surface soil and inhalation of vapors while working in the upland portion of the Site. The occupational worker scenario assumes that workers do not frequent the beach portion of the Site during typical work activities.
 - O Upland Construction/Excavation Workers. This scenario relates to workers performing utility upgrades or maintenance or other activities that involve the disturbance of soil at the Former Gas Works Property and the properties in the vicinity. Upland construction workers could be exposed to Site-related COCs through dermal contact with and incidental ingestion of soils and inhalation of vapors. Typical construction worker activities (e.g., grading or excavation for building foundations) are expected to extend up to approximately 3 feet in depth.²³
 - O Potential Future Residential Users of the Site (Not a Current or Planned Use). The Former Gas Works Property and the properties in the vicinity are zoned for industrial uses, and this is expected to remain the case for the foreseeable future. However, the potential for exposures of future residents may be appropriate to evaluate as part of the risk assessment to understand potential implications should properties within the Site be converted to residential uses. On-site residents could be exposed to Site-related COCs

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²³ For the purposes of the RI, "surface soil" is defined as the 0- to 3-foot interval, based on likely ecological and human health exposure scenarios.

through direct contact with and incidental ingestion of surface soils and inhalation of vapors. No water supply wells are located on or near the Former Gas Works Property, but consumption of groundwater has been retained as a potential pathway for screening, pending further evaluation of groundwater beneficial uses.

The Site and vicinity are used by a variety of upland and aquatic species. An initial list of species common to the region has been compiled (Table 4-1), using locally available published sources. Listed in the table are species that use or may occasionally use the Site and vicinity. The species listed in Table 4-1 are grouped into representative categories to illustrate different ecological exposure pathways.

Preliminary representative receptors that could potentially use the Site and be directly or indirectly exposed to contaminated media were identified from the list of species common to the region (Table 4-1). EPA guidance is available to help identify receptors potentially at risk (EPA, 1992, 1997a, and 1998b). Receptors potentially at risk include:

- Federal or state rare, threatened, or endangered species
- Resident species or communities with the greatest exposure to chemicals in sediment and surface water
- Species or functional groups that are essential to, or indicative of, the normal functioning of the affected habitat

Based on review of the USFWS catalogue of federally listed species that could potentially be within the Site vicinity (2016), the following five species are listed:

- Bull trout (Salvelinus confluentus)
- Killer whale (Orcinus orca)
- Marbled murrelet (Brachyramphus marmoratus)
- Streaked horned lark (Eremophila alpestris strigata)
- Yellow-billed cuckoo (Coccyzus americanus)

WDFW also identifies species with a State endangered, threatened, sensitive, candidate or monitored status (WDFW, 2016). Regionally common species with a Federal or Washington State listing are identified in Table 4-1.

At the Site, the ecological receptors potentially at risk include the animals and plants that use the terrestrial and/or aquatic habitats within the Site. These were categorized into plant, invertebrate, reptile and amphibian, fish and shellfish, and bird and mammal groups. Species within these groups likely to be present at the Site were determined through review of relevant regional information. The representative species from these groups were selected as target species to be evaluated in the risk assessment. A brief description of each group is as follows.

Plants. Both terrestrial and aquatic plants are important resources because they provide significant habitat for fish and wildlife. Terrestrial vegetation is limited in the Site and primarily exists along the bluff. The riparian zone is primarily forested riparian and is generally present across the Site shoreline. Aquatic vegetation in Dyes Inlet is

patchy. Dyes Inlet and Sinclair Inlet do not support any floating kelp and non-floating kelp species are present in just 18% of the shoreline throughout the entire basin (Redman et al., 2005).

Invertebrates. The invertebrate community (both terrestrial and aquatic) is an important receptor group because soil and benthic invertebrates consume plants and detritus, provide critical nutrient cycling, and represent a trophic link to other organisms that consume them. Because invertebrates, with the exception of crabs, are relatively sessile and are in direct contact with soil or sediment, they provide an integrated measure of toxicity. Benthic organisms observed in the intertidal and nearshore of Sinclair inlet include crustaceans, molluscs, arthropods, polychaetes and echinoderms (GeoEngineers, 2011; KiTSA, 2012).

Reptiles and Amphibians. Reptiles potentially using the Site include common garter snakes (Thamnophis spp.). Frogs, turtles, salamander and newts inhabit the Sinclair Inlet Watershed (KiTSA, 2012), but are not likely to be present at the Site due to unsuitable habitat. Reptiles and amphibians will not be directly assessed in the ERA because exposure models and toxicological data for reptiles and amphibians are limited. There is significant uncertainty associated with the exposure and sensitivity of this group of receptors to COPCs at the Site. Given the limited information available, a meaningful assessment of risk to this group of receptors is not possible. However, it is assumed that the risk characterization and risk-based management for other assessment endpoints (e.g., soil invertebrates, mammals, fish, and birds) will provide protection of reptiles and amphibians at the Site.

Fish and Shellfish. Fish and shellfish are key elements of freshwater, estuarine, and marine ecosystems for a number of reasons. As one of the most diverse groups of vertebrates, fish are able to occupy a wide range of ecological niches and habitats. As such, fish represent important components of aquatic food webs by processing energy from aquatic plants (i.e., primary producers), zooplankton and benthic macroinvertebrate species (i.e., primary consumers), or detrivores. Fish also represent important prey species for piscivorous wildlife including reptiles, birds, and mammals. Fish species present in Sinclair and Dyes Inlets includes representatives of the benthivorous, omnivorous and piscivorous guilds. Common species are summarized in Table 4-1.

Birds. Birds likely to be found at the Site are grouped as primarily marine-dependent or terrestrial-predators. The primary terrestrial avian feeding guild at the Site is avian predators which includes robins and crows. Aquatic-dependent species guilds include piscivorous raptors and other shorebirds. Species common to Sinclair and Dyes Inlets include great blue heron, osprey (KiTSA, 2012), and sediment-probing birds such as the sandpiper (Buchanan, 2006).

Mammals. Mammals common to the region and likely to be present at the Site include occasional coyotes, rabbits, squirrels, voles, shrews, mice, moles, and raccoons (KiTSA, 2012). Mammals will be grouped into terrestrial and aquatic-dependent receptors for the ERA. Primary terrestrial mammal feeding guilds include herbivorous mammals, insectivorous mammals, omnivorous mammals, and carnivorous mammals. Aquatic-dependent mammals common to the region and likely to be present at the Site for extended periods of time comprise members of the piscivorous feeding guild including Dall's porpoise, harbor porpoise, California sea lion, harbor seal, northern sea lion, and

river otter (KiTSA, 2012). Other marine mammals including several species of whales have been occasionally sighted near the Site, but at much lower frequencies.

Representative preliminary receptors for the ERA were selected based on the following criteria:

- Ecological relevance;
- Potential levels of exposure to COPCs;
- Social or economic importance;
- Sensitivity to COPCs; and
- Availability of sufficient natural history information to allow meaningful assessment of exposure and risk.

Exposure pathways relevant to these representative species are presented in Figure 4-4 for aquatic (i.e., fish) and aquatic-dependent (e.g., heron and river otter) receptors and in Figure 4-5 for terrestrial receptors.

Figure 4-4 shows the aquatic wildlife receptors with potentially complete exposure pathways: direct contact with and ingestion of sediment, porewater, and marine water; and consumption of benthic invertebrates, fish, and other potentially contaminated prey. The representative aquatic receptors listed in Figure 4-4 include the following:

- Piscivorous Mammals (e.g., Harbor Seals). There is a potential for limited exposure of piscivorous mammals foraging at the Site. The harbor seal was selected to represent mammals with primarily aquatic diets feeding mostly on demersal and pelagic fish with some crustaceans and mollusks. Potentially complete exposures are associated primarily with consumption of aquatic biota and, to a lesser extent, with direct contact with and ingestion of sediment and marine surface water.
- **Piscivorous Raptors (e.g., Ospreys).** There is a potential for limited exposure of piscivorous raptors foraging at the Site. The osprey represents birds that feed primarily on pelagic fish with some demersal fish. Potentially complete exposures are associated primarily with consumption of aquatic biota and, to a lesser extent, with direct contact with and ingestion of marine surface water.
- Shore Birds (e.g., Herons and Sandpipers). There is a potential for exposure of shore birds residing or foraging at the Site. The great blue heron represents birds that feed primarily on demersal fish with some pelagic fish and crustaceans. The spotted sandpiper represents shore birds that obtains much of their diet by probing or "mining" soft sediments along shorelines and that feed on aquatic worms, mollusks, and crustaceans. Potentially complete exposures are associated primarily with consumption of aquatic biota, direct contact and incidental ingestion of sediment, and, to a lesser extent, with direct contact with and ingestion of marine surface water.
- **Piscivorous Fishes (e.g., Rockfish).** Piscivorous fishes feed on higher trophic level species and may be at increased risk from bioaccumulative COCs. Non-listed rockfish represent fish residing or foraging at the Site that may potentially be

- exposed to Site-related COCs primarily through consumption of fish and direct contact with marine surface water and sediment, incidental ingestion of marine surface water, and to a lesser extent, ingestion of sediment.
- Omnivorous Fishes (e.g., Sculpins). Sculpin represent omnivorous fishes residing or foraging at the Site that may potentially be exposed to Site-related COCs primarily through consumption of aquatic biota, direct contact with sediments and marine surface water, incidental ingestion of marine surface water, and to a lesser extent, ingestion of sediment and consumption of other biota.
- Benthivorous Fishes/Shellfish (e.g., Flatfish, Bivalves, and Crabs).

 Benthivorous fish/shellfish prey on infaunal and epibenthic organisms. English sole or other flatfish, and crabs represent benthivorous fishes/shellfish residing or foraging at the Site that may potentially be exposed to Site-related COCs primarily through consumption of biota and through direct contact with and ingestion of sediments and marine surface water, and to a lesser extent through consumption of other biota.
- Benthic Invertebrates (e.g., Benthic Infauna Community). Benthic invertebrates residing at the Site may potentially be exposed to Site-related COCs through direct contact with and ingestion of sediments, porewater and marine surface water, and, to a lesser extent, through consumption of other biota.
- Macrophytes (e.g., Algae and Kelp). Macrophytes residing at the Site may potentially be exposed to Site-related COCS through direct contact with sediment and porewater and marine surface water.

The upland properties at the Site have historically been developed and used for industrial operations. However, portions of these properties include habitat that could be used by terrestrial ecological receptors. These areas primarily include the vegetated areas of the Former Ravine and the bank. Terrestrial ecological receptors with potentially complete exposure pathways are illustrated on Figure 4-5 and include the following:

- Avian Predators (e.g., Robins). There is a potential for exposure of avian predators foraging or nesting at the Site. The robin represents birds that prey on soil invertebrates and, to a lesser degree, fruit. The primary exposure pathways include the consumption of terrestrial biota and direct contact with and incidental ingestion of Site soil, and to a lesser extent, contact with and ingestion of on-Site surface water.
- Carnivores (e.g., Coyotes). There is a potential for limited exposure of carnivores foraging at the Site. The coyote represents upper-trophic level mammals that prey primarily on small mammals and soil invertebrates. The primary exposure pathways for these receptors include the consumption of terrestrial biota and direct contact with and incidental ingestion of Site soil, and, to a lesser extent, contact with and ingestion of on-Site surface water.
- Omnivores (e.g., Raccoons). There is a potential for limited exposure of omnivores foraging at the Site. The raccoon is a common, medium-sized, opportunistic feeder with a varied diet depending on season and location; the raccoon was selected to represent mammals feeding primarily on soil invertebrates and plants. The primary exposure pathways for these receptors include the

- consumption of terrestrial biota and direct contact with and incidental ingestion of Site soil and, to a lesser extent, contact with and ingestion of on-Site surface water.
- **Herbivores** (e.g., Voles). There is a potential for exposure of herbivores residing at the Site. The vole is a common small mammal that consumes shoots, grasses, and bark and is prey for carnivorous mammals and birds. The vole represents mammals feeding solely on plants. The primary exposure pathways for these receptors include the consumption of terrestrial biota, direct contact with and incidental ingestion of Site soil, and, to a lesser extent, contact with and ingestion of on-Site surface water.
- Insectivores (e.g., Shrews). There is a potential for exposure of insectivores residing on the Site. The shrew represents mammals feeding primarily on earthworms and other soil invertebrates. The primary exposure pathways for these receptors include the consumption of terrestrial biota, direct contact with and incidental ingestion of Site soil, and, to a lesser extent, contact with and ingestion of on-Site surface water.
- **Upland Vegetation.** There is a potential that plants growing at the Site could be exposed to Site-related COCs in soil. The primary exposure pathways for plants is direct contact with Site-related COCs in the soil.
- Soil Invertebrates. There is a potential for exposure of earthworms and other biota living at the Site. The primary exposure pathways for these receptors include direct contact and incidental ingestion of Site-related COCs in soil, consumption of terrestrial biota, and, to a lesser extent, contact with and ingestion of on-Site surface water.

4.4 Preliminary Contaminants of Potential Concern

This Section identifies preliminary COPCs based on: (1) contaminants typically associated with the former gas works process (carbureted water gas); (2) contaminants associated with other potential historical sources within the initial study area (ISA; see Section 5.1); (3) contaminants detected during previous Site investigations; and (4) other EPA contaminants of interest. The COPCs and (following completion of the baseline risk assessments) ultimately the COCs, that are determined to apply to the Site-related decisions may include some, none, or all of the contaminants identified in this Section. The COCs that are ultimately determined to apply to the Site-related decisions will be established in the baseline risk assessments on the basis of data and information that are collected as part of the RI/FS process (Section 5.2.4).

Contaminants typically associated with carbureted water-gas manufacturing processes include the following:

- Light aromatic hydrocarbons, such as BTEX compounds;
- Heavier aromatic hydrocarbons, including PAHs;
- Other SVOCs, such as tar acids (e.g., phenol and cresols) and heterocyclic aromatics (e.g., carbazole and dibenzofuran); and
- Cyanide and sulfides associated with spent purifier materials.

Other historical processes with the potential for releases at the Site include petroleum transfer and storage, metal fabrication, and vehicle and equipment salvage and repair. Contaminants typically associated with these processes include solvents (VOCs), petroleum hydrocarbons (including BTEX and PAHs), and metals.

Though there are no existing data indicating their presence at elevated concentrations, EPA has identified polychlorinated biphenyls (PCBs), pesticides and dioxins/furans as other contaminants of interest at the Site. PCBs are man-made organic chemicals, manufactured between 1929 and 1979, and used in industrial and commercial applications including electrical, heat transfer, and hydraulic equipment; in paints, plastics and rubber products; and in pigments and dyes. PCBs may still be present in products and materials that were manufactured before 1979, including electrical transformers and capacitors, fluorescent light ballasts, adhesives, oil-based paint and caulking. Pesticides are substances, or mixtures of substances, intended for preventing, destroying, repelling, or mitigating any living organisms (e.g. insects, mice, weeds, fungi, microorganisms) that occur where they are not wanted or that cause damage to crops, humans or other animals. The term pesticides applies to insecticides, herbicides, fungicides, and various other substances used to control pests. Dioxins/furans are a group of toxic chemical compounds that are created unintentionally as a result of human activities, such as production of herbicides, combustion processes (waste incineration or burning wood, coal or oil fuel), and chlorine bleaching of pulp and paper, as well as natural processes like forest fires (http://www.epa.gov/dioxin/learn-about-dioxin).

The preliminary COPCs for the Site fall within the following groups of contaminants:

- VOCs, as identified and quantified by EPA Method 8260C.
- SVOCs, including carcinogenic- and non-carcinogenic PAHs, as identified and quantified by EPA Method 8270D/SIM.
- Cyanide, as identified and quantified by EPA Method 9014.
- Metals, as identified and quantified by EPA Methods 200.8/6010/6020/7471B.
- PCBs, as identified and quantified by EPA Method 8082.
- Pesticides, as identified and quantified by EPA Method 8081B.
- Dioxins/furans, as identified and quantified by EPA Method 1613.

Table 4-3 identifies the specific contaminants within each group that are considered preliminary Site COPCs. Non-toxic metals including calcium, chloride, iodine, magnesium, phosphorous, potassium and sodium, are essential nutrients and are not identified as COPCs even though some of them have been previously detected at the Site. The preliminary COPCs were selected if information indicates they are confirmed or suspected to be present at the Site.

Table 4-3 is not intended to provide an exhaustive and complete list of all COPCs for the Site. The scope of work for the RI will include analysis and reporting of the full standard list of contaminants for each analytical method, as described in detail in the Upland SQAPP and Marine SQAPP (Appendices A and B, respectively). Initial Site investigations, which will investigate and characterize potential sources of contamination at the Site (see Section 5.5), will generally include analysis of all samples for the methods

listed in the bullets above²⁴. The data collected during this first phase of work will be screened against initial PRGs. The baseline risk assessments (to be prepared in parallel with the RI Report) will determine which of these COPCs that presents an unacceptable risk are to be identified as COCs, based on the data gathered during the RI. COCs identified in the baseline risk assessments will then be carried forward into the FS for the evaluation of remedial options.

[.]

²⁴ As described in Section 5.5, certain analyses may be excluded if field observations indicate high levels of contaminants (e.g., tar) that would cause analytical interferences for other compounds (e.g., pesticides), and a tiered approach to selecting samples for dioxin/furan analysis will be implemented.

5 Work Plan Rationale

This Section describes the basis and approach for the RI data collection program. It includes the following information:

- Description and basis for the initial study area (ISA) that is to be characterized during the RI (Section 5.1);
- Summary of data needed to complete the RI and FS (Section 5.2);
- Approach for completing the risk assessment (Section 5.3);
- DQOs for collected data (Section 5.4);
- The approach for filling data gaps (Section 5.5); and
- Potential contingency studies that may be required after initial data collection has been completed (Section 5.6).

Details of the specific sampling and analysis programs for the upland and marine areas are provided in the Upland and Marine SQAPPs (Appendices A and B).

5.1 Initial Study Area

The purpose of defining the ISA is to provide a focused area for sampling and analysis in the initial phase of the RI/FS (AOC, EPA, 2013a). The ISA is not intended to define the Site boundaries. The Statement of Work (SOW) for the AOC anticipates "the ISA will encompass the area of operation of a former manufactured gas plant (MGP)..., including the area where contaminants from the area of operation have come to be located, which includes upland, beach and sediments." The ISA has been developed according to the guidelines established by the SOW and includes an upland portion and a sediment portion. The rationale for the upland and sediment portions of the ISA is explained further in the following Sections.

5.1.1 Upland Initial Study Area

The upland portion of the ISA (Figure 5-1) includes the Former Gas Works Property and portions of neighboring properties where gas works operations, including byproduct storage and disposal, are documented or suspected to have occurred. This includes the northern portion of the Penn Plaza Property where a drip tank was located and the eastern portion of the Sesko Property where materials from the former gas works process may have been placed in the Former Ravine. The upland portion of the ISA also includes areas where contamination associated with operations other than the former gas works could potentially be commingled with contamination from the gas works. These non-gas-works operations include the former Lent's bulk petroleum storage tank farm on the Sesko Property, petroleum pipelines located in the northern portion of the Penn Plaza Property and the Sesko Property, and various light industrial operations on the McConkey and Penn Plaza Properties.

Consistent with the SOW, the proposed ISA encompasses all upland areas where contaminants associated with the former gas works are likely to be located. The existing data collected from areas near the boundaries of the ISA suggest that contamination associated with the former gas works may not extend beyond the ISA. More data are needed to determine if this is the case. The existing data include the results of soil and groundwater sampling from well MW-1 on the Penn Plaza Property, borings MP03 and MP02 within Thompson Drive, borings SP01 and SP02 on the Sesko Property, and explorations associated with the Former SC Fuels Property to the east of the ISA.

The first phase of the RI will characterize the nature and extent of contamination within the ISA and assess the subsurface characteristics that may influence the migration of contaminants. These data will be used to determine where additional investigation may be warranted. Investigations outside of the ISA, if needed, would then be specifically designed and implemented to focus on the characterization of identified issues.

5.1.2 Sediment Initial Study Area

The sediment portion of the proposed ISA (Figure 5-2) comprises intertidal and subtidal areas in the general vicinity of the Former Gas Works Property. The sediment portion of the ISA is described as follows:

- Historical potential source areas associated with the former gas works (including the Former Gas Works Dock and the former drainage line) have been included.
- All beach sediments adjacent to the Former Gas Works Property that exhibited elevated PAH concentrations during the 2013 TCRA have been included.
- The offshore boundary of the ISA extends out past midchannel in the Port Washington Narrows, well past the bathymetric low point in the channel. This addresses potential migration pathways associated with groundwater and/or NAPL migration and those associated with potential sediment transport.
- The eastern and western boundaries of the ISA extend between 500 and 1,000 feet in an east-west direction from the Former Gas Works Property, allowing documentation of the potential transport of sediments that may have resulted from the east-west tidal currents within the Port Washington Narrows.

The sediment portion of the ISA includes multiple potential sources that are unassociated with historical activities on the Former Gas Works Property: multiple historical petroleum transfer docks, multiple stormwater and CSO outfalls, and the Port Washington Marina.

As part of the RI/FS activities related to sediments, there is a need to understand regional sediment quality or water quality that may affect either current Site conditions or result in future recontamination of the Site. Therefore, sampling activities for sediments and surface water will not be exclusively confined to the ISA. Some sampling during the RI/FS will occur outside the sediment portion of the ISA. However, the investigation and remediation of non-Site-related contaminant sources that are located outside the ISA is not an objective of this RI/FS.

5.2 Data Needs

The data needs have been identified through the RI/FS scoping process and development of the Scoping Memorandum (Aspect and Anchor QEA, 2015). This Section discusses the data needs that affect all components of the RI/FS process. The general data needs, specific data gaps, and planned RI data collection methods for the upland and marine portions of the Site are summarized in Tables 5-1 and 5-2, respectively. The general approach for addressing the data needs is summarized in Section 5.5.

5.2.1 Upland Data Needs

5.2.1.1 Site Physical Characteristics

Characterization of the physical properties of the soil is necessary to evaluate the contaminant migration pathways and the remedial options. Soil samples will be collected from all typical lithologic units, as feasible, for physical characterization to include grain size, density, moisture content, and organic carbon content.

The data needs associated with the hydrogeology of the Site include data to define aquifer and aquitard units across the Site, evaluate the hydraulic conductivity of aquifer units, and understand the influence of tidally influenced surface water on groundwater flow and contaminant transport from the Site. The installation and sampling of groundwater monitoring wells is needed to provide these physical data, as well as samples to define the extent of groundwater contamination. The distribution of groundwater contaminants is associated with groundwater flow, which may be affected by seasonal variations in groundwater levels due to precipitation, as well as interaction with surface water. The information needed to satisfy these data needs will be obtained by sampling groundwater for chemical and geochemical parameters, logging geologic information, measuring static and transient water levels, and performing aquifer testing.

5.2.1.2 Nature and Extent of Contamination

A primary objective of the RI is to delineate the nature and distribution of contamination in the potentially affected media at the Site, which include soil, groundwater, air, surface water, and sediment. Samples of each potentially affected medium will be collected for chemical analysis of the Site COPCs throughout the RI process.

Also important in understanding the nature and extent of contamination is to identify and delineate contaminant source materials such as NAPL. As described in Section 3.7.1, NAPL has been observed in Site soil. Based on historical Site use, both LNAPL (density is less than that of water) and DNAPL (density is greater than that of water) may be present. If there is sufficient volume and the soil is sufficiently permeable, both LNAPL and DNAPL will migrate downward via gravity flow through the soil. Because it is less dense than water, LNAPL will begin to migrate laterally when it encounters groundwater, primarily in the direction of groundwater flow. DNAPL is denser than water and will continue to sink below the water table. As it migrates downward, both in the vadose zone and through the water-bearing zone, NAPL leaves behind a residual coating of product on the soil grains, which can be used as an indicator of the potential presence of NAPL.

DNAPL will continue to migrate downward via gravity flow until the available volume of mobile DNAPL has been depleted or until a soil layer with lower permeability is encountered. DNAPL may collect in pools on top of low-permeability layers and migrate laterally through seams of higher permeability soil. Downward vertical migration of DNAPL below the water table can also be slowed or eliminated by an upward hydraulic gradient. Along with the evaluation of the presence of NAPL, the geologic and hydrogeologic conditions at the Site will be characterized as part of the evaluation of potential NAPL mobility.

Because NAPL is a hazardous substance, but also a potential source of contaminants to other media, the characterization of the presence, nature, and extent of NAPL will be another primary objective of the RI. The data needs associated with NAPL include investigation to identify its presence, collection of data to delineate its lateral and vertical extent in the subsurface, and laboratory testing to evaluate its composition and mobility. The information needed to satisfy these data needs will be obtained by field screening soil, gauging monitoring wells for the presence of NAPL, evaluating chemical data from soil, groundwater, and sediment for indications of NAPL presence²⁵, and, if feasible, collecting NAPL samples for physical and chemical testing.

5.2.1.3 Contaminant Fate and Transport

Contaminants present in Site media may migrate from one location to another via the fluid flow processes of advection or diffusion, transfer between media via partitioning mechanisms, and attenuate as the result of physical, chemical, or biological processes. Contaminants can also be transformed into different chemicals or destroyed by biological or chemical reactions. Understanding contaminant migration and transformation across the Site is important for evaluating potential exposure pathways, anticipating how the nature and extent of contamination may change over time, and evaluating the potential effectiveness of remedial actions, including estimating the restoration time frame. The potential contaminant migration pathways and transformation processes are described in detail in Section 4.2.

To evaluate fate and transport of upland contaminants, it will be necessary to collect data to evaluate potential medium-to-medium migration pathways and NAPL migration pathways (Table 5-1). The data needs associated with the evaluation of upland contaminant fate and transport include data to define the physical characteristics of soil and NAPL, define the physical characteristics of aquifers and aquitards, evaluate natural attenuation and degradation of contaminants in soil and groundwater, and evaluate groundwater chemical data to assess spatial and temporal trends. Information obtained to determine the physical characteristics of the Site (Section 5.2.1) and the nature and extent of contamination (Section 5.2.2) will be used to evaluate contaminant fate and transport. The additional information needed to satisfy these data needs will be obtained by the collection and analysis of groundwater samples for specific indicators of natural

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²⁵ Concentrations of hydrocarbons in soil greater than 10,000 mg/kg generally indicate the potential presence of tar or NAPL (Cohen and Mercer, 1993). The detection of benzene, naphthalene, or PAHs in groundwater at a concentration greater than 10 percent of each contaminants' solubility suggests that NAPL may be present at or upgradient of that location.

attenuation or degradation of contaminants, including geochemical indicators of contaminant degradation and the evaluation of groundwater data for changes in contaminant concentrations along a chemical flow path or over time.

Data collected to inform the evaluation of contaminant transport within and between environmental media and evaluate potential mechanisms for contaminant attenuation include:

- Physical soil characteristics, including soil type, grain size, density, and TOC
 content, to support the analysis of migration pathways including the potential for
 contaminants to leach from soil into groundwater and to sorb to soil from
 groundwater.
- Hydraulic characteristics, including hydraulic conductivity, groundwater gradients, and tidal influences, to evaluate groundwater flow and associated contaminant transport.
- Groundwater geochemical data, including dissolved organic carbon, nitrate, nitrite, sulfate, sulfide, ferrous iron, dissolved manganese, alkalinity, pH, dissolved oxygen and oxidation-reduction potential, to evaluate natural attenuation and biological and chemical degradation of contaminants.

Geochemical data and the change in chemical concentrations along groundwater flow paths may be used for qualitative evaluations of the occurrence of contaminant degradation or for qualitative evaluations in conjunction with contaminant fate-and-transport groundwater models.

Temporal data collected at a Site well network will be used to assess seasonal trends (e.g., by looking at fluctuations during quarterly sampling events) in contaminant transport and longer-term trends to assess whether contaminant plumes are stable, expanding, or shrinking.

Soil and groundwater chemical data, along with physical characteristics, will be used to evaluate potential migration pathways to soil vapor and indoor air. The potential for vapor intrusion will be assessed in accordance with guidance (EPA, 2015) and may include vapor intrusion modeling (e.g., the Johnson-Ettinger model) to assess potential impacts under current or future uses. Potential additional soil vapor or indoor air studies are discussed in Section 5.6.

5.2.1.4 Risk Assessment

The data needs for the risk assessment generally overlap those for the RI and FS. Specific types of information required to support the development of a baseline human health risk assessment (HHRA) and a baseline ERA for the upland areas include the following:

- Conduct supplemental testing within the upland portion of the ISA to finalize the list of COPCs for the upland area.
- Determine the nature and extent of contamination in surface soil and subsurface soil to assess risks for human and ecological receptors.

- Develop sufficient data to estimate potential risks related to the effect of contaminant vapor on indoor air quality, including shallow subsurface soil and/or groundwater quality data or soil vapor data.
- Determine the nature and extent of contamination in groundwater to assess risks for human and ecological receptors.

5.2.2 Marine Data Needs

5.2.2.1 Site Physical Characteristics

To evaluate physical forces and overall geologic formations in the sediment portion of the ISA and the adjacent portions of the Port Washington Narrows, evaluations of current velocity, and sediment substrate studies by means of a towed video camera are needed. Current velocity will be measured at two depth profiles (near-bottom and mid-channel) along each transect and will be used to indicate potential impacts of current velocity on sediment stability within the ISA and the Port Washington Narrows. Sediment grain size results will be used in conjunction with modeled wind/wave and measured tidal current velocities to evaluate intertidal and subtidal sediment transport processes in the Port Washington Narrows.

Similarly, towed-camera surveys will be conducted to document the sediment substrate type, natural, and anthropogenic features in perpendicular and parallel transects in the vicinity of the sediment ISA and the adjacent Port Washington Narrows.

5.2.2.2 Nature and Extent of Contamination

A primary objective of the RI is to delineate the nature and lateral/vertical distributions of contamination in the surface water and sediment. Samples of each potentially affected medium will be collected for chemical analysis of the Site COPCs²⁶ throughout the RI process.

Because NAPL is a hazardous substance, but also a potential source of contaminants to other media, the characterization of the presence, nature, and extent of NAPL will be another primary objective of the RI. The data needs associated with NAPL include investigation to identify its presence, collection of data to delineate its lateral and vertical extent in the subsurface, and laboratory testing to evaluate its composition and mobility.

5.2.2.3 Contaminant Fate and Transport

To evaluate fate and transport of marine contaminants, it will be necessary to collect data to evaluate medium-to-medium migration pathways and NAPL migration pathways. These data needs will be satisfied by an evaluation of surface sediments, surface sediment porewater, subsurface sediments, surface water, and physical characteristics of sediments.

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²⁶ Except a tiered approach will be used to select samples for analysis of VOCs and dioxins/furans. Sediment samples will be analyzed for VOCs based on field screening as described in Appendix B. Sediment and surface water samples will be archived for possible analysis of dioxins/furans, following the protocol described in Section 5.5.2.

In addition, data are needed to characterize the physical mechanisms of transport within the Port Washington Narrows to determine potential transport through surface water, sediment littoral drift, and sediment bed load mobility.

5.2.2.4 Risk Assessment

The data needs for the risk assessment generally overlap those for the RI and FS. Specific types of information required to support the development of a baseline human health risk assessment (HHRA) and a baseline ERA for the marine areas include the following:

- Conduct supplemental testing within historical source areas to confirm the list of COPCs for the marine investigation.
- Determine the nature and extent of Site-associated PAH contamination in surface sediments.
- Evaluate potential PAH contamination in surface water within the marine portion of the ISA.
- Determine the nature and extent of Site-associated PAH contamination in subsurface sediments in the beach area for use in evaluating potential risks for beach construction workers.
- Assess the partitioning behavior of PAHs in surface sediment using solid-phase microextraction (SPME) to inform whether literature-based partitioning estimates provide a reasonable basis for estimating bioavailability, or whether additional data are needed to perform the risk assessment.
- Where warranted following initial data collection and in consultation with EPA and the Tribe, implement contingent bioassay testing and/or sediment porewater testing to augment sediment and porewater data and evaluate potential impacts on benthic infaunal communities.
- Where warranted following initial data collection and in consultation with EPA
 and the Tribe, implement contingent tissue testing of selected species to refine
 estimates of potential bioaccumulation of contaminants in aquatic species that are
 harvested by seafood consumers or that serve as prey for higher trophic level
 ecological receptors.
- Use video surveys to augment available literature regarding the aquatic species that may use the Site and vicinity.
- Use beach surveys to assess the current abundance of shellfish potentially subject to harvest activities in beach areas near the Site.

Section 5.3 describes how each of the data collection activities will be used in support of the risk assessment activities.

5.2.3 COC Identification

The scope of work for the RI/FS will include collection and analysis of samples for Site COPCs and contaminants of interest (see Section 4.4) to support the identification of Site COCs, which are those contaminants identified to be present at concentrations that pose a potential risk to human health or the environment in media for which there is a potential complete exposure pathway. The final COCs will be defined based on the results of the baseline risk assessments (Section 5.3). The basis for eliminating a contaminant or contaminant group as a COPC include the following:

- The contaminant is a naturally occurring compound and is detected within the range of a documented natural background concentration.
- The contaminant is not identified as a COC in the baseline human health or ecological risk assessments (see Section 5.3).

5.3 Risk Assessment Approach and Methodology

Consistent with the AOC, a baseline ERA and HHRA will be performed to support the RI/FS decision-making. The baseline risk assessments will be completed in parallel with the preparation of the Draft RI Report. While the initial phase of investigation is being completed, ongoing consultation with the EPA and the Tribe will be conducted to identify seafood consumption based PRGs. The seafood consumption-based PRGs will be included in the Risk Assessment Technical Memorandum. The Risk Assessment Technical Memorandum will identity any data gaps and the means by which they should be addressed (e.g., seafood tissue sampling, additional soil, or sediment testing).

The data collection activities associated with the risk assessment will be conducted as part of the Site characterization activities. The planned data collection activities will address the data needs for completion of the risk assessment for all receptors and exposure scenarios identified in Section 4.3.

The specific risk assessment plan for the HHRA is presented in Table 5-3. The risk assessment plan for the baseline ERA is presented in Tables 5-4 and 5-5. The tables provide the following information:

- The receptor to be evaluated;
- The evaluation framework to be used to estimate potential risks for that receptor under the specific exposure scenario;
- The RI data that will be used in support of the risk assessment for the specific exposure scenario; and
- The endpoint and interpretive framework to be used to quantify potential risks.

5.3.1 Risk Assessment Technical Memorandum

An interim deliverable, the Risk Assessment Technical Memorandum, will be used to document the preliminary screening of the collected RI data and provide a detailed description of the methods to be used for the baseline risk assessments. The Risk

Assessment Technical Memorandum will also identify data gaps and define the scope of additional data collection, where necessary. The Risk Assessment Technical Memorandum will define the scope and methods for any tissue testing required to complete the Baseline Risk Assessments. The Risk Assessment Technical Memorandum will be prepared in conjunction with the Phase 1 Data Report, ²⁷ which is discussed further in Section 6.3. Consultation between the EPA and the Tribe will inform the development of seafood consumption-based PRGs.

As described in Section 4.4, the preliminary COPCs were selected based on: (1) contaminants typically associated with the former gas works process (carbureted water gas); (2) contaminants associated with other potential historical sources within the initial study area (ISA; see Section 5.1); (3) contaminants detected during previous Site investigations; and (4) other EPA contaminants of interest. The Risk Assessment Technical Memorandum will re-evaluate the EPA-identified contaminants of interest (i.e., PCBs, dioxins/furans, and pesticides) to determine if they will be retained as COPCs in the ERA or HHRA.

The Risk Assessment Technical Memorandum will provide the following information identified in Tables 5-3, 5-4, and 5-5:

- The specific data to be used for the evaluation of each exposure scenario;
- Results of preliminary data screenings;
- Statistical approaches (where applicable) to be used to estimate exposure point concentrations for each exposure scenario;
- Description of any models or calculations to be used to estimate exposures, including the following:
 - Methods used to estimate soil vapor and indoor air quality from soil and groundwater data;
 - Source of any biota-sediment accumulation factors to be used to estimate the bioaccumulation of sediment contaminants in aquatic species;
 - Partitioning coefficient values used to estimate porewater quality from bulk sediment data;
 - Models and parameters used to estimate the total daily intake of contaminants for each receptor; and
 - The rationale for seafood consumption rates used for fish and shellfish consumption exposure scenarios, as developed in consultation with EPA and the Tribe.
- Applicable toxicity information and exposure parameters; and

²⁷ In the AOC, this report is also called the RI/FS Data Report.

• Current screening levels, benchmarks, and toxicity reference values to be used in the ERA and HHRA.

The Risk Assessment Technical Memorandum will also identify data gaps and contingent testing activities (where applicable) to be implemented in support of the risk assessment. The data gap identification will include the environmental media (e.g., soil, sediment, tissue) and COPC/COPC group where data are insufficient to prepare the ERA or HHRA. Any proposed testing activities will be documented in an addendum to this Work Plan in accordance with the AOC (see Section 6.2).

5.3.2 Human Health Risk Assessment

The HHRA methodology will be based on national and regional guidance designated by EPA, including, but not limited to, the following:

- Risk Assessment Guidance for Superfund (RAGS), Volume I: Human Health Evaluation Manual, Parts A through F (EPA, 1989);
- Interim Guidance: Developing Risk Based Clean-up Levels at Resource Conservation and Recovery Act Sites in Region 10 (EPA, 1998a);
- Risk Assessment Guidance for Superfund, Volume I: Human Health Evaluation Manual (Part E, Supplemental Guidance for Dermal Risk Assessment, Final) (EPA, 2004);
- Risk Assessment Guidance for Superfund, Volume I: Human Health Evaluation Manual (Part F, Supplemental Guidance for Inhalation Risk Assessment) (EPA, 2009);
- Human Health Evaluation Manual, Supplemental Guidance: Update of Standard Default Exposure Factors (EPA, 2014);
- The Exposure Factors Handbook (EPA, 2011); and
- The Framework for Selecting and Using Tribal Fish and Shellfish Consumption Rates for Risk-Based Decision Making at CERCLA and RCRA Cleanup Sites in Puget Sound and the Strait of Georgia (EPA, 2007).

Toxicity data will be developed on the basis of the EPA hierarchy of human health toxicity values (EPA, 2003). Any updates to the above sources will be documented in the Risk Assessment Technical Memorandum.

The Draft Baseline HHRA Report will be submitted to EPA 180 days after the receipt of validated data from samples collected during the Site characterization activities. The Final Baseline HHRA Report will be included in the Final RI Report.

5.3.3 Ecological Risk Assessment

The ERA methodology will address both terrestrial and aquatic ecological exposures for the receptors identified in Section 4.3. The ERA methodology will be based on EPA guidance, including, but not limited to, the following:

- Ecological Risk Assessment for Superfund: Process for Designing and Conducting Ecological Risk Assessments, (EPA, 1997a)
- Guidelines for Ecological Risk Assessment (EPA, 1998b); relevant and appropriate updated EPA guidance material (e.g., EPA's Eco Updates)
- EPA Region 10 Supplemental Ecological Risk Assessment Guidance for Superfund (EPA, 1997b).

Toxicity data will be developed in accordance with EPA guidance (e.g., EcoSSLs) and databases (e.g., Ecotox), peer-reviewed scientific literature, and recent EPA-approved risk assessments. Any updates to the above sources will be documented in the Risk Assessment Technical Memorandum.

The Draft Baseline ERA Report will be submitted to EPA 180 days after the receipt of validated data from samples collected during the Site characterization activities. The Final Baseline ERA Report will be included in the Final RI Report.

5.4 Data Quality Objectives

The data needs for the RI/FS have been developed through a methodical planning process to ensure appropriate sampling, analyses, and data evaluations are conducted to meet the RI/FS objectives. EPA's Guidance on Systematic Planning Using the Data Quality Objective Process (EPA, 2006) was used to acquire the necessary data to develop the RI/FS, through a process called Data Quality Objectives (DQO). The seven-step DQO process is a tool to determine the type, quantity, and quality of data necessary for any subject analysis. The seven steps are:

- 1. State the problem;
- 2. Identify the Goal of the Study;
- 3. Identify Information Inputs;
- 4. Define the Boundaries of the Study;
- 5. Develop the Analytic Approach;
- 6. Specify Performance or Acceptance Criteria; and
- 7. Develop the Plan for Obtaining the Data.

DQOs were developed to address the data needs summarized in Tables 5-1 and 5-2. The following 10 DQOs have been identified to complete the RI/FS:

- 1. Contamination in Soil and Groundwater (Table 5-6)
- 2. Source of Contamination (Upland) (Table 5-7)

- 3. Site Physical Characteristics (Table 5-8)
- 4. NAPL Characterization (Table 5-9)
- 5. Contaminant Fate and Transport (Table 5-10)
- 6. Habitat and Intertidal Shellfish Surveys (Table 5-11)
- 7. Contamination of Surface Sediment (Table 5-12)
- 8. Contamination of Subsurface Sediment (Table 5-13)
- 9. Contamination of Surface Water (Table 5-14)
- 10. Marine Area Sediment Stability and Recontamination Processes (Table 5-15)

These DQOs are summarized in Tables 5-6 through 5-15, and are further detailed in Section 5.5. The characterization approach to fulfill these DQOs is described in the following Sections.

5.5 Remedial Investigation Approach

This Section presents the general approach for characterizing the Site and addressing data gaps related to the upland and marine portions of the Site. The CSM will be continuously updated as data is collected and evaluated to modify sampling locations and approaches to meet the objectives of each phase of the investigation. Specific sampling and analysis details including preliminary exploration locations, sampling and analysis rationale, and field procedures are provided in the Upland SQAPP (Appendix A) and Marine SQAPP (Appendix B).

5.5.1 Upland Investigation

The upland investigation will be conducted sequentially to adaptively manage the scope of work to address specific objectives. The investigation will start with evaluating potential sources and work its way out from those sources to determine the nature and extent of contamination at the Site. The upland RI activities will consist of five general phases of work, as follows:

- 1. Geophysical Investigation.
- 2. Source Characterization.
- 3. Source Areas Investigation.
- 4. Outside Source Areas Investigation.
- 5. Groundwater Testing and Monitoring.

Data gathered during each step of the field investigation will be used to guide subsequent data collection.

A detailed scope of work, including objectives, approach, and proposed explorations, for performance of the Geophysical Investigation and the Source Characterization is provided in the following sections. These two components will be sequenced so that the results of

the Geophysical Investigation can be reviewed before the Source Characterization to determine whether additions or modifications to the scope of work are warranted.

The objectives and approach for the Source Areas Investigation and the Outside Source Areas Investigation are also discussed in the following sections. These include a rationale for determining exploration methods, locations, and other details, such as depth and sampling intervals. Some details, such as the number, location, and depth of explorations, will be determined based on the results of the preceding steps.

This Work Plan is designed to provide sufficient detail to enable decision-making by the project team in the field, to streamline the investigation program. Key decision points or unanticipated conditions that would require consultation with EPA for resolution are identified. A field communication plan to ensure that field activities and data are communicated to the Cascade and EPA project teams in a timely manner is included in Appendix A and summarized in Section 9.2.

Depending on the results of the upland investigation, contingent investigations or studies that would require an addendum or addenda to this Work Plan may be warranted. Some potential contingent investigations are described in Section 5.6. The process for planning and reporting on additional phases of investigation work is described in Section 6.2.

5.5.1.1 Geophysical Investigation

The Geophysical Investigation will consist of geophysical and utility surveys to provide preliminary information regarding subsurface conditions. The primary objective of the Geophysical Investigation is to evaluate the former gas works operations area and the Former Ravine for potential buried structures (i.e., piping, tanks and equipment foundations) or anomalous ground conditions that may indicate historical use of the subsurface (i.e., covered and filled pits) or fill material. The surveys will also be used to identify active storm drain lines or other existing utilities. The results of the surveys will be used to identify potential sources for further assessment in the Source Characterization, and to determine if adjustments to the Source Characterization, such as moving or adding explorations to investigate observed anomalies, are warranted.

The geophysical surveys, including the field data collection and interpretation, will be completed by hydroGEOPHYSICS, Inc. Three separate geophysical surveys will be conducted successively, as described below, to meet different objectives in different areas of the Site.

5.5.1.1.1 Electromagnetic Induction (EM) Survey.

The objective of the EM survey is to identify subsurface metallic objects, variations in soil moisture content that may indicate shallow subsurface filled pits or trenches, and to identify areas where shallow subsurface fluid or contaminant has pooled. Electromagnetic field data are collected using portable ground conductivity instrumentation. A transmitting coil induces an electromagnetic field and a receiving coil at a fixed separate distance measures the amplitudes of the in-phase and quadrature components of the electromagnetic field. The in-phase component is most sensitive to metallic objects. The quadrature (also referred to as conductivity) component is sensitive to soil condition variations. High density EM data covering large areas can be collected within a short time period and readily processed and visualized for quick interpretation. Depending on

subsurface conditions, EM data can evaluate conditions up to 15 feet below ground surface.

The EM survey area includes the entirety of the upland portion of the ISA except for the Shoreline fill area, where the character of the slope prevents safe data collection, and the nearshore section of the marine portion of the ISA, as depicted on Figure 5-3. Electromagnetic induction data will be acquired along parallel survey lines over the survey area at a line spacing of approximately 3 feet, varied as necessary based on accessibility. EM data will not be collected where buildings or immobile metallic objects are located at the ground surface because of potential electromagnetic interference. The EM data is expected to be collected within one day and can typically be processed and visualized overnight to allow for an evaluation of the findings and refinement of the approach for the ground penetrating radar (GPR) survey.

5.5.1.1.2 Ground Penetrating Radar Survey

The GPR survey will provide targeted, high resolution characterization of the location, size and dimensions of subsurface metallic objects on the McConkey Property identified during the EM survey. In addition, the GPR survey will evaluate the McConkey Property for non-metallic subsurface infrastructure and variations in subsurface conditions that may be indicative of covered and filled pits or limits of fill material. The GPR survey will only be completed on the McConkey Property because of uneven or poorly accessible ground surface conditions on other portions of the Site that prevent GPR equipment mobilization.

GPR data will be collected along orthogonally arranged lines at a distance of approximately 6-feet over areas of interest identified in the EM survey, with a maximum survey area of the entire paved or bare and accessible portions of the McConkey Property (Figure 5-3). The data will be collected using a Sensors and Software Noggin 250 Smartcard, which includes a 250 MHz shielded antenna, with a maximum depth of penetration of 8 to 10 feet depending on soil conductivity. GPR data will be filtered and processed by the geophysical consultant in the office using software to reduce noise within the data while drawing out GPR targets and identifying the location and depth of any GPR targets detected.

5.5.1.1.3 Electrical Resistivity Survey

The electrical resistivity (ER) survey will be used to provide high resolution, 2-dimentional cross-sections through the subsurface of the Former Ravine to determine the lateral limits and depth of fill material. The ER technique is assumed to be well suited to meet this objective based on the anticipated contrast in electrical properties between the fill material and the native lithology.

The ER survey will be performed using a SuperstingTM R8 multichannel electrical resistivity system and 18-inch long electrodes, installed 8- to 10- inches into the ground, on 2-foot spacing along three transects across the Former Ravine (Figure 5-3). This will provide high resolution imaging and an expected depth of investigation of approximately 30 feet.

5.5.1.2 Source Characterization

The objective of the Source Characterization is to identify, delineate and characterize sources at the Site. A 'source' is defined as media that exhibits gross contamination such as tar or NAPL; materials coated by, or saturated with, NAPL; or other MGP-related

feedstocks or byproducts such as ash, slag, or purifier waste. Field observations that may indicate source material are identified in Appendix A.

The specific objectives of the Source Characterization are to:

- Identify and delineate subsurface features (such as tanks, sumps, and piping) that may be associated with sources;
- Identify and evaluate sources at the Site;
- Delineate the lateral and vertical extent of sources; and
- Identify the COPCs associated with each source.

The scope of work for the Source Characterization has been developed based on what is currently known about the Site and its operational history, and does not include any planned explorations on the eastern side of the Sesko Property or around the southwest corner of the upland ISA because there are no known or suspected historical sources located in these areas²⁸. However, if the geophysical surveys identify any subsurface anomalies warranting investigation in these areas, additional explorations will be added to the Source Characterization. Results of the Geophysical Investigation and proposed modifications to the Source Characterization will be reviewed with EPA prior to conducting the Source Characterization.

This section is organized as follows:

- Characterization Methods and Process
- Exploration Locations
- Sample Collection and Analysis

5.5.1.2.1 Source Characterization Methods and Process

In shallow soils, test pits and trenches are likely to be more effective than borings at evaluating the presence and characteristics of sources because of the ability to make more extensive observations. Direct-push probes will be used in areas where test pits and trenches are impracticable (e.g., beneath or adjacent²⁹ to structures) and may be used to vertically delineate sources at depths beyond what is achievable with test pits or trenches. However, because of the density of native glacial soils beneath the former gas works operations area and the suspected presence of buried debris in the Former Ravine, the practical depth of direct-push soil borings at the Site is expected to also be limited to relatively shallow soils. Deeper borings, if needed to characterize or delineate sources, would be advanced using hollow-stem auger or sonic drilling methods³⁰.

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²⁸ There is the potential for unknown sources to be present that may not be identified by geophysical methods. The Outside Source Areas Investigation (Section 5.5.1.4) will provide additional exploration of areas outside of known source areas.

²⁹ Trenches generally will not be excavated at locations and depths that would result in greater than a 1.5:1 H:V slope from building foundations.

³⁰ Source Characterization activities will be scheduled and sequenced to minimize delays in mobilizing additional equipment to the Site if needed. For instance, a sonic drill rig may be contingently scheduled in advance to follow direct-push borings. If a direct-push boring reaches refusal, Source

The process for identifying, characterizing, and delineating suspected sources using trenches is as follows:

- 1. Excavate a trench to observe subsurface conditions in the vicinity of the suspected source. The trenches will be aligned to provide the highest probability of encountering a source, if present, based on available historical information.
- 2. If a subsurface feature representing a potential source (e.g., sump, tank or pipe) is observed, the extent of the feature will be determined through excavation to identify the limits of the feature as practicable³¹.

3. If a source is observed:

- a. The trench will be extended to determine the lateral extent and dimensions of the source as practicable, in the same manner as the investigation into the extent of a subsurface feature described above. Where trenching cannot define the extent of a source, other methods of investigation will be used to meet the objectives.
- b. At the location where the source is strongest (i.e., more extensive or more highly contaminated), a cross-trench (approximately perpendicular to the original trench) will be excavated to determine the lateral extent and dimensions of the source.
- c. Once identified and delineated laterally, the vertical extent of each source or source area will be determined by advancing at least two soil borings immediately adjacent to the source/source area³².
- 4. The source will be considered delineated when field observations no longer indicate gross visible contamination, strong odors, or significantly elevated PID readings, as determined and documented by the field lead.

At locations or depths where trenches are not feasible, direct-push borings with continuous core soil samples will be used to determine the presence of source material and delineate the lateral and vertical extent of sources. If source material is observed at a boring, stepout borings in opposite directions at 20-foot intervals will be advanced. Similar to

Characterization explorations (i.e., trenches or direct-push borings) in other areas of the Site may be advanced pending mobilization of larger equipment.

³¹ Shallow historic/abandoned piping may be excavated and removed, if practicable and necessary to meet the objectives of the Source Characterization. The removal of piping will be completed through excavation and removal from the ground to a practicable extent, which may correspond to subsurface limitations (e.g. a building foundation), an aboveground structure, property boundaries beyond which excavation work may require street use permits or approvals of adjacent property owners, a depth beyond which an exploration excavation is no longer feasible without structural support or shoring, or other practicable limits. The pipe will be capped or plugged at the practicable limits of exploration and removal

³²These source delineation borings will not be advanced through the source material to minimize potential for carry down of contaminants on drilling equipment. If it is necessary to drill through the source material, because of access or other logistical limitations, appropriate drilling methods and precautions (see Appendix A) will be employed to minimize potential carry down.

trenches, a cross-transect of borings at 20-foot intervals will be advanced at the location where the source is strongest, based on field observations.

If explorations onto adjacent street rights-of-way or properties are needed, approval by the property owners (e.g., via an access agreement or street-use permit) will be sought. If further investigation into piping location is warranted beyond practicable excavation limits, other methods may be employed to meet the investigation objectives (i.e. utility location, GPR surveys, etc.). If piping remains in place beyond the feasible extent of removal, the end will be capped and sealed, and the GPS coordinates of its location will be recorded for future reference. If the origin of the piping remains unclear at the limits of feasible removal, a camera survey or further geophysical survey may be conducted in an attempt to identify its origin and historic use.

Source Characterization explorations will be completed through fill materials and into native soils, where practicable. Soil borings completed to delineate the vertical extent of each source or source area will be advanced to 16 feet below ground surface or to 6 feet below the source material, whichever is deeper, unless refusal is met shallower. If refusal is encountered within fill material (e.g., on buried debris) before the target depth is reached, the boring will be relocated within five feet of the previous location. If refusal is met a second time, the exploration location will be abandoned and alternative investigation methods will be evaluated. If refusal is encountered due to dense native soils before the target depth is reached, alternative drilling methodologies (e.g., hollow-stem auger or sonic) will be employed to reach the target depth.

Trenches will be completed to native soil or a minimum depth of 6 feet bgs, unless health and safety considerations (e.g., sidewall sloughing) dictate otherwise. The maximum depths of test pits and trenches will be determined by the field lead based on the source being investigated, observations made during the investigation, the subsurface lithology and the limitations of the equipment given the location and surface conditions of the exploration. As examples:

- Explorations intended to evaluate potential releases from surface sources, such as
 aboveground tanks used to store oil, gasoline or finished gas, will be completed at 6
 feet bgs if no indications of releases are observed. If source material is observed in the
 exploration, the exploration will be completed to define the vertical extent of source
 material or the maximum depth that is safely feasible, whichever is shallower.
- Explorations intended to evaluate potential releases from shallow subsurface sources, such as underground piping or structures such as tar wells, will be completed at depths sufficient to determine the depth of the structure and the conditions beneath the structure, and to a minimum 6 feet bgs if no indications of releases are observed.
- Explorations to evaluate the gas holder as a suspected subsurface source area will be completed at depths sufficient to determine the depth of the gasholder and the conditions at the base of the gasholder, or the maximum depth that is safely feasible, whichever is shallower.
- Explorations intended to evaluate the character of fill material in the Former Ravine will be completed to the shallower of native soil or the maximum safe depth of exploration, if surface conditions/ground stability will not safely support construction equipment large enough to extend the exploration deeper.

5.5.1.2.2 Exploration Locations

Proposed exploration locations, based on current information, are shown on Figure 5-4. These include evaluation of former gas works features and fill areas, as follows:

• Former Gas Works Features. Former gas works features, such as the gas holder, process equipment, and feedstock and byproduct storage areas, are a logical place to start the Source Characterization as those features may have resulted in releases, deposition or burial of source material. The locations of the former gas works features and the Source Characterization explorations to evaluate them are shown on Figure 5-4. Some of these features are visible today. The location of the former gas holder is evident as a circular outline in the asphalt. Likewise, there is an expression in the asphalt in the approximate location of the former scrubber. Additionally, a portion of the concrete slab where the coal/coke briquettes were stored is still present and visible at the ground surface. The locations of other former gas works features and other potential source areas will be estimated using field global positioning system (GPS) equipment based on their approximate coordinates obtained from georeferenced historical aerial photographs.

Trenches are impractical in the vicinity of the former tar pit because of a building overlying this area (Figure 5-4). Five soil borings will be advanced inside of the building to evaluate the former tar pit. If a source is observed in any one of the borings by the field lead, a step-out exploration will be completed at a distance of approximately 20 feet from the original boring. Step-out explorations outside of the building may be completed as either another boring or as an excavated trench, based on the target depth and field conditions.

• Ravine and Shoreline Fill Areas. Sources may also be present in/as fill material in the ravine or along the shoreline. The Source Characterization will include excavation of trenches in these areas to evaluate the nature and extent of fill material, including confirming the extent of fill material, as practicable given safety limitations, and to identify and characterize sources. Preliminary trench locations for the Source Characterization are depicted on Figure 5-4.

Manhole A is currently filled with concrete debris and dirt, which is unlikely to be successfully removed without the removal of the manhole structure itself. Therefore, the Source Characterization will include the removal of Manhole A through excavation and temporary shoring, observation of surrounding soils, and a camera survey of any inlets identified.

The results of the Geophysical Investigation may indicate subsurface anomalies, lateral and vertical limits of fill material, and/or buried metallic objects outside of the current proposed exploration plan. Source Characterization explorations may be modified or added based on the results of the Geophysical Investigation as follows:

- Explorations will be completed in locations where a buried metallic object is identified through the EM and GPR surveys.
- Explorations will be completed in locations where EM and GPR survey results identify subsurface anomalies on the McConkey Property that may indicate filled pits, concrete structures, pooled fluids or buried debris.

5.5.1.2.3 Sample Collection and Analysis

Soils collected from borings, test pits, and trenches will be characterized by soil type and field screened for indications of COPC impacts and NAPL presence (as discussed in more detail in the Upland SQAPP [Appendix A] and Section 5.5.1.2.3), and the results will be recorded. Continuous soil samples will be collected from soil borings for logging and field screening.

To evaluate the COPCs associated with identified sources, samples of source materials (e.g., NAPL or NAPL-coated soil) will be collected for laboratory analysis of the COPCs³³. At least one representative sample of each distinct source material will be collected for chemical characterization. If sufficient free-phase NAPL can be collected, NAPL samples will also be collected and submitted for petrophysical testing (density, viscosity, flashpoint).

As described above, borings will extend to a minimum of 6 feet below observed sources. To evaluate the vertical distribution of COPCs in soil beneath sources, samples from soil borings advanced to delineate source depth will be collected for COPC analysis³⁴ from the following depth intervals:

- Less than 2 feet below the source; and
- 2 to 6 feet below the source.

If borings extend more than 6 feet below the source, additional samples will be collected at 4-foot intervals and archived. Archived samples will be analyzed for COPCs that exceed PRGs in the interval above the archived sample.

5.5.1.3 Source Areas Investigation

After the Source Characterization is complete, the Source Areas Investigation will be conducted to evaluate the nature and extent of contamination attributable to identified sources or areas where discrete sources may be contributing to a larger, general area of source material when similar source materials are located in relatively close proximity (Source Areas). Source Areas will be defined based on the characteristics and location of the various sources identified during the Source Characterization. This section describes the objectives, logic, and process of conducting the Source Areas Investigation. Identification of Source Areas and specific exploration locations and target depths will be proposed and reviewed with EPA prior to conducting this investigation.

The objectives of the Source Areas Investigation are as follows:

- To characterize the magnitude and vertical extent of soil and groundwater contamination within and downgradient of each Source Area.
- To assess the presence of free-phase NAPL in the saturated zone and evaluate NAPL characteristics if present.

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³³ Except a tiered approach will be used to select soil samples for analysis of dioxins/furans, following the protocol described in Section 5.5.1.3.1.

³⁴ Selection of soil samples within a targeted depth interval is described in Appendix A.

This investigation will include advancement of soil borings for characterization of subsurface lithology; collection of soil samples for laboratory analysis; and installation of monitoring wells for gauging NAPL and collecting groundwater samples for laboratory analysis. The exploration locations will include the following:

- Locations downgradient of Source Areas identified during the Source Characterization, particularly those with potentially mobile contaminants such as NAPL, to determine the lateral and vertical extent of contamination, including NAPL, in soil.
- Along groundwater flow transects, including locations along the shoreline, to evaluate
 potential migration of contaminants to the Port Washington Narrows and to identify
 appropriate locations and construction details for the installation of groundwater
 monitoring wells.

The Upland SQAPP (Appendix A) provides the sampling and analysis details to meet these objectives, including soil classification, field screening, and soil and groundwater sample collection. The process for conducting the investigation, including decision criteria for identifying exploration locations and details, is described below.

5.5.1.3.1 Source Areas Investigation Process

Source Area explorations will proceed in the following sequence:

- Completing a transect of deep borings oriented perpendicular to groundwater flow immediately downgradient³⁵ of each Source Area, to determine the lateral and vertical extent of soil contamination and identify appropriate monitoring well locations.
- Completing one or more wells along each transect based on boring results.
- Completing one or more wells within each Source Area.

The location and depth of deep borings in the transect will be determined following the decision logic described in Appendix A. Approximately four borings will be advanced per transect; however, the exact number of borings will depend on the size of the Source Area and the proximity of adjacent Source Area transects.

These borings will be advanced using Sonic drilling technology, which enables collection of continuous core for lithologic description and field screening and has the highest likelihood of reaching intended depths. At least one soil sample will be collected from these borings from each of the following units: fill material, native soils in the vadose zone, and each saturated unit and aquitard encountered during the boring. Each sample will be analyzed for all Site COPCs except dioxins/furans. Samples for potential dioxin/furan analysis will be archived, and archives samples will be analyzed for dioxins/furans if PCBs, chlorinated phenols, or chlorinated pesticides are detected.

³⁵ It is assumed that the downgradient flow direction is north, toward the Port Washington Narrows. This assumption will be re-evaluated during early hydrogeologic characterization (see Section 5.5.1.4).

At least one monitoring well will be constructed based on field screening and analytical data³⁶. The monitoring well will target the area of highest contamination observed along each transect. If contamination is detected in soil or through field observations in more than one hydrogeologic unit, an additional well will be installed in each potentially impacted unit.

After the installation and development of new monitoring wells, and redevelopment and evaluation of existing wells to ensure that they are of satisfactory integrity for use, groundwater samples will be collected for chemical analysis to evaluate the lateral and vertical distribution of COPCs³⁷ in groundwater from all existing and viable wells, including those installed previously by others. Groundwater samples will also be analyzed for conventional and geochemical parameters³⁸. At the time of sampling, all new and existing viable wells will be evaluated for the potential presence of NAPL If NAPL is observed, the thickness of NAPL will be measured. If sufficient NAPL volume is present to collect a NAPL sample, a sample from each well containing NAPL will be collected and analyzed for petrophysical parameters.

5.5.1.4 Outside Source Areas Investigation

The investigation activities to be conducted as part of the Outside Source Areas Characterization are those primarily intended to characterize the physical site environment to support development of the conceptual site model (i.e., to define the characteristics of soil and groundwater to support fate and transport evaluations), to bound extent of contamination and identify Site boundaries, and to complete the risk assessment.

Specific objectives of the Outside Source Areas Investigation are as follows:

- Evaluate the characteristics of native soil;
- Identify and characterize water-bearing zones and aquitards;
- Conduct a preliminary evaluation of groundwater flow direction to support well placement;
- Characterize shallow soil conditions to bound Site contamination and support the risk assessment; and
- Characterize groundwater quality cross-and up-gradient of source areas to define the lateral extent of COPCs in groundwater.

Some of these objectives may be met through the collection of data or information during the Source Characterization (Section 5.5.1.2) and the Source Areas Investigation (Section

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³⁶ Work will be scheduled and sequenced to minimize downtime between borings and well installation. Borings for each Source Area transect will be completed first. Drill rigs to install monitoring wells will be scheduled in advance to coincide shortly after receipt of analytical data. Pre-validated data will generally be considered adequate for decision making. Analysis of lab results may be expedited if needed to prevent significant delays to field activities.

³⁷Except groundwater samples will be archived and analyzed for dioxins/furans only if PCBs, chlorinated phenols, or chlorinated pesticides are detected in the sample.

³⁸ Conventional and geochemical parameters include dissolved organic carbon, nitrate, nitrite, sulfate, sulfide, ferrous iron, dissolved manganese, alkalinity, pH, dissolved oxygen, oxidation-reduction potential, and other major ions (sodium, potassium, chloride, calcium, magnesium).

5.5.1.3). For example, samples of fill material and native soil will be collected from borings advanced as part of the Source Areas Investigation.

The Outside Source Areas Investigation will be conducted after the Source Areas Investigation is complete and Source Areas have been defined. However, preliminary deep borings outside the area where Source Areas are anticipated will be completed prior to the Source Areas Investigation to provide a preliminary aid to that investigation.

The Outside Source Areas Investigation will include the following components:

- Preliminary deep soil borings and wells to evaluate Site physical characteristics.
- Soil sampling using Incremental Sampling Methodology (ISM) to evaluate the nature and extent of soil contamination outside Source Areas.
- Perimeter well installation to characterize the nature and extent of groundwater contamination outside Source Areas.

These components are described below.

5.5.1.4.1 Preliminary Deep Borings and Wells

This work will consist of completing approximately three deep borings to evaluate and physically characterize Site lithology, including distinct geologic units, water-bearing zones and aquitards. Approximate locations are shown on Figure 5-4. The total depth of the borings will be dependent on observed geologic and hydrogeologic conditions at the time of drilling; the rationale for the total depth of the preliminary deep borings is presented in detail in Appendix A. Borings will be advanced using Sonic drilling methods to collect continuous soil samples for lithology and achieve target depths. Soil samples from each distinct lithologic unit will be analyzed for soil physical and conventional properties, including grain size, total organic carbon, and Atterberg limits. Soil density will be determined using SPT.

These borings will be completed as wells either deeper in the water table aquifer or in an underlying aquifer following the decision steps in Appendix A. The purpose of these wells is to characterize hydrogeologic characteristics of deeper groundwater. A tidal study will be conducted using existing shallow wells and the new deep wells to provide a preliminary estimate of groundwater flow direction and gradients. This work will be done prior to the Source Areas Investigation, as the data will be used to help determine the location and completion details of Source Area borings and wells. Salinity will be measured at these wells as part of evaluation of hydrogeologic conditions, but these wells will not be sampled for COPCs unless determined appropriate later in the investigation process.

5.5.1.4.2 Shallow Soil Characterization

Shallow soils outside Source Areas will be characterized using incremental-sampling methodology (ISM). ISM is a structured composite sampling and processing protocol having specific elements designed to reduce data variability and increase sample representativeness for a specified volume of soil under investigation (ITRC, 2012). ISM may provide a reasonable approach to collect the data necessary to calculate mean contaminant concentrations to assess risks to human and ecological receptors in shallow soil. In ISM, the area to be characterized is divided into distinct Decision Units based on historical operations, expected types or magnitudes of contamination, and/or potential

exposure scenarios. ISM is generally not appropriate to characterize areas containing sources. Because of the large density of sampling points, ISM may also have limited applicability in areas with significant utility constraints, such as street rights-of-way. Decision Units will be determined based on the results of the Source Characterization and Source Areas Investigation. The application and appropriateness of ISM at the Site, including its ability to meet investigation objectives, may be reassessed after initial investigations are completed.

Details of ISM sampling are described in Appendix A. ISM sampling will include sampling of two depth intervals: 1) surface soil (0 to 3-foot depth); and 2) shallow subsurface (3- to 6-foot depth) soil. These intervals are based on consideration of human health and ecological exposure pathways for the risk assessment. The ISM process involves the following:

- Collecting a large number of soil samples from each Decision Unit and depth interval spatially distributed across the Decision Unit. For this Work Plan, three triplicates of 30 samples per Decision Unit and depth horizon (i.e., 180 samples per Decision Unit) are assumed.
- Compositing each set of 30 samples and analyzing each composite sample for all COPCs except VOCs³⁹ and dioxin/furans⁴⁰ (i.e., 6 samples per Decision Unit).

ISM results in the outermost Decision Units (i.e., at the ISA boundary) will be compared to PRGs. If PRGs are exceeded, step-out Decision Units may be identified based on the results, or an alternative sampling plan to bound the extent of contamination in Site soil (i.e., discrete sampling at soil borings) may be identified.

5.5.1.4.3 Cross- and Up-Gradient Groundwater Quality Characterization.

Monitoring wells will be installed to provide groundwater quality data in cross- and upgradient locations from the Source Areas and to define the Site boundary. Wells will be developed and sampled for all COPCs⁴¹ and for dissolved organic carbon, nitrate, nitrite, sulfate, sulfide, ferrous iron, dissolved manganese, alkalinity, pH, dissolved oxygen, oxidation-reduction potential, and other major ions (sodium, potassium, chloride, calcium, magnesium) to characterize Site geochemical conditions.

5.5.1.5 Groundwater Testing and Monitoring

After Site boundaries are identified, a groundwater testing and monitoring program will be implemented to more fully characterize groundwater conditions. The testing and monitoring program will include:

 Slug testing at selected monitoring wells to measure hydraulic conductivity of each water-bearing zone.

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³⁹ Samples for VOC analysis should not be composited. One discrete VOC sample from each depth horizon will be collected for every 250 square feet of Decision Unit area.

⁴⁰ Except soil samples will be archived and analyzed for dioxins/furans only if PCBs, chlorinated phenols, or chlorinated pesticides are detected in the sample.

⁴¹ Except groundwater samples will be archived and analyzed for dioxins/furans only if PCBs, chlorinated phenols, or chlorinated pesticides are detected in the sample.

- A tidal study using pressure transduces in Site wells to measure groundwater gradients, and tidal influences.
- Quarterly measurement of groundwater conventional and geochemical parameters, including dissolved organic carbon, nitrate, nitrite, sulfate, sulfide, ferrous iron, dissolved manganese, alkalinity, pH, dissolved oxygen, oxidation-reduction potential, and other major ions (sodium, potassium, chloride, calcium, magnesium) to evaluate conditions for biological and chemical degradation of contaminants.
- Quarterly measurement of selected COPCs for a minimum of one year, to assess seasonal trends.

Wells for hydraulic testing will be determined after the Site boundaries and the number of affected water-bearing zones have been determined.

As described in previous sections, wells will be tested initially for COPCs and conventional/geochemical parameters. Specific wells and analyses for quarterly groundwater monitoring will be determined based on initial testing results. Locations and frequency for groundwater monitoring after one year, to evaluate long-term trends, will be determined based on the first years' worth of data.

5.5.2 Marine Investigation

The marine investigation elements described in this Section address data gaps identified in the RI/FS scoping process. The locations of each data acquisition point/location have been determined in consultation with EPA. However, each of the proposed locations is subject to revision based on conditions encountered in the field at the time of data acquisition.

The sequence of the investigation elements is integrated into this Work Plan to inform PRG development, COPC identification, risk assessment, and the definition of the nature and extent of contamination. Sediments and surface water will be analyzed for Site COPCs (see section 4.4) with the exception of VOCs and dioxins/furans. Sediments will be field screened for VOCs and selected for analysis based on protocols described in the Marine SQAPP (Appendix B). Sediment and surface water samples will be archived and selected for analysis of dioxins/furans only where PCBs, chlorinated phenols, or chlorinated pesticides are detected in the samples. The elements of the marine investigation are summarized in Table 5-16.

The initial data acquisition program will include the following:

- **Video Surveys.** Video surveys will be conducted to identify substrate, habitat characteristics, and presence/abundance of aquatic resources near the Site. This information will be used to evaluate the presence of anthropogenic structures or features that require consideration during the RI/FS (Figure 5-5).
- **Tidal Current Evaluation.** Near-bottom and mid-depth tidal currents within the aquatic areas of the Site will be monitored to assist in the evaluation of sediment stability and sediment transport processes (using sediment grain size testing results)
- **Surface Sediment Investigation**. Surface sediments will be sampled and analyzed as follows:

- Within the marine environment immediately adjacent and to the north of the Former Gas Works Property, 17 surface samples will be collected and analyzed for Site COCPs (Figure 5-6).
- Within the ISA, 16 surface samples will be collected and analyzed for Site COPCs (Figure 5-7).
- Outside of the ISA, additional data will be acquired to assess the physical characteristics of sediment within the Port Washington Narrows (Figure 5-8). These data will be used to evaluate sediment transport processes within the Port Washington Narrows.
- At 5 intertidal locations with the ISA, paired samples of bulk sediment and porewater will be analyzed to evaluate Site PAH as compared to literature-derived partitioning coefficients (Figure 5-7).
- **Baseline Shellfish Habitat Surveys.** Beach surveys will be performed to evaluate the distribution and abundance of existing shellfish species within and near the beach areas adjacent to the Former Gas Works Property (Figure 5-7 and Figure 5-8). These habitat surveys will provide information on existing habitat conditions. The findings of the surveys are not intended to restrict shellfish consumption rate estimates used in the Human Health Risk Assessment.
- Subsurface Sediment Investigation. Subsurface sediment core samples will be collected from the beach and subtidal areas sloping down into the Port Washington Narrows to evaluate the vertical distribution of chemicals, including the potential presence of NAPL and hydrocarbon sheen, in subsurface sediments.
- Surface Water Investigation. Surface water samples from selected Site and background locations will be collected and analyzed during multiple sampling events to assess potential variability in the concentrations of contaminants in surface water (Figure 5-7 and Figure 5-8). The surface water analysis will include a full suite of conventional and Site COPC testing.

The general rationale and approach for these components of the marine investigation are described in the following Sections, and the details are included in Appendix B. Based on the results of the marine investigation, contingent investigations or studies may be warranted; those are described in Section 5.6.

5.5.2.1 Video Survey

The objective of the towed camera surveys is to identify substrate types, habitat characteristics, the presence/abundance of aquatic resources, and identify any unknown anthropogenic features (e.g., outfalls, structures, or sunken barges). The video surveys will be collected along 12 predefined transects in the Port Washington Narrows in the vicinity of the ISA (Figure 5-5). Six transects each will be conducted perpendicular to and parallel with the shoreline of the Port Washington Narrows. The parallel video transects are positioned at the southern and northern shores at the -10 feet mean lower low water (MLLW) and -20 feet MLLW contours (Figure 5-5), through the deeper channel area adjacent to the former gas works, and over the shallower area in the central channel. One of the perpendicular transects is positioned through the slope adjacent to the former gas works and two are positioned to the east and west in the Port Washington Narrows. After

the video surveys are complete, the locations of the transects will be plotted on a figure. The videos will be reviewed to qualitatively determine, at a minimum, the substrate type, habitat characteristics, presence/abundance of aquatic resources, and any other significant observations, and the results will be logged. This survey will yield an interpretative figure that presents the video survey findings. The results will be used to determine habitat types and identify anthropogenic structures or natural features or resources that merit consideration as part of the RI/FS process.

5.5.2.2 Tidal Current Evaluation

Tidal current surveys will be conducted by a qualified contractor along four transects at the locations shown on Figure 5-5. A vessel-mounted acoustic Doppler current profiler will be used to measure current velocity along transects over the course of a daily tide cycle with a relatively high tidal exchange. Sampling will be performed during a period of high tidal exchange (between a high tide of at least mean higher high water and a low tide below MLLW). Measurements will be collected in both directions (i.e., back and forth) across each transect location to decrease any directional bias in the data. Results from near-bottom measurements within the ISA will be used to inform the FS and assess the potential impacts of tidal currents on sediment stability.

5.5.2.3 Beach Shellfish Surveys

Beach shellfish surveys will be conducted to document the types and quantities of species currently present within and immediately adjacent to the ISA. The surveys will be conducted at seven locations within and adjacent to the ISA and (Figure 5-6 and Figure 5-7) in accordance with WDFW methods (Appendix B; Campbell, 1996). These surveys will provide information on existing habitat conditions. The findings of the surveys are not intended to restrict shellfish consumption rate estimates used in the Human Health Risk Assessment.

5.5.2.4 Surface Sediment Investigation

Surface sediment samples will be collected to characterize the lateral nature and extent of contamination, evaluate chemical fate and transport, determine COPCs, evaluate relative bioavailability of PAHs, and quantify sediment transport processes. All surface sediment samples will be collected from a depth 0 to 4 inches below the mudline that typically constitutes the bioactive zone. Consistent with previous Site-related investigations, ISA intertidal sediment samples will be collected by hand during low tide. All subtidal surface sediment samples and intertidal samples outside the ISA will be collected using a power actuated Van Veen grab sampler. The surface sediment samples submitted for differing testing suites are described as follows:

- To characterize the lateral nature and extent of contamination, and evaluate chemical fate and transport, 19 surface sediment samples will be collected and analyzed, for Site COPCs, immediately adjacent and to the north of the former gas works (Figure 5-6). At these locations the analytical suite includes alkylated PAH analysis. The alkylated PAH results will be used to assess the differentiation of potential PAH sources detected within the sediments.
- To determine the relative bioavailability of PAHs, five intertidal locations will be tested by ex situ SPME methodology (Figure 5-6). The methodology includes bulk

sediment collection, insertion into the sediments of performance reference compound-spiked SPME fibers, equilibration period, post-equilibration SPME chemical analysis, and data evaluation to derive estimated porewater concentrations. Paired with bulk surface sediment results, these data will be used to evaluate bioavailability relative to literature-based values. Ex situ testing methodology is further detailed in Appendix B.

• To define the lateral extent of contamination in sediments within the ISA, 16 additional surface sediment locations will be collected (Figure 5-8). These samples will be analyzed for Site COPCs as detailed in Appendix B.

To evaluate sediment transport processes, 16 samples outside the ISA but within intertidal and subtidal bedded sediment will undergo physical testing. Physical testing coupled with current velocities and modeled wind and wave action will identify conditions that mobilize and redistribute sediment. Nineteen surface sediment sampling locations immediately adjacent to the former gas works, within the marine portion of the ISA, were selected in consultation with EPA (Figure 5-6). 17 of those sampling locations are arrayed in transects downslope toward the Port Washington Narrows channel. Within the marina to the west of the slope, two additional sampling locations were placed to evaluate sediment quality where historical dredging has been conducted. These 19 sampling locations are collocated with subsurface cores collection locations for vertical delineation of the nature and extent of contamination (see 5.5.2.5).

Sixteen additional surface sediment locations will be sampled to characterize the lateral nature and extent of contamination in areas of the ISA further offshore of the former gas works. For representative spatial coverage emanating from the former gas works, a sample will be collected from the western extent of the marina, from two intertidal locations in the eastern extent of the ISA, from four subtidal locations immediately offshore of the base of the slope, and from seven subtidal locations distributed throughout the ISA.

Surface sediment locations outside the ISA will be submitted for physical testing to inform evaluations of sediment transport processes (coupled with ADCP results and modeled wind and wave conditions) within Port Washington Narrows (Figure 5-8). The sediment transport in intertidal areas (through littoral drift) and bedded sediments (channel sediment stations) will be evaluated. Of the 16 locations, 11 target the littoral drift zones and five characterize bedded sediments in the channel of Port Washington Narrows.

The results of all aspects of the surface sediment investigation will be presented in the Phase 1 Data Report.

5.5.2.5 Subsurface Sediment Investigation

Subsurface core sampling will be conducted to determine the vertical nature and extent of Site-related COPCs (including NAPL and sheen). The subsurface explorations will be advanced at 17 sampling locations along transects aligned down the slope from the Former Gas Works Property and at two locations immediately west of the slope within the marina, all of which are at the same locations as the surface sediment sampling locations (Figure 5-6). The subsurface sampling area includes the intertidal areas where with Site-related COPCs are known to be elevated and in locations of historical dock structures. As designed, the core sampling program is of sufficient density to evaluate migration

pathways to characterize all potential directions of travel as described in Section 4.2.1. To evaluate potential release pathways to the Port Washington Narrows, the deepest core in each transect targets the -20 feet MLLW elevation to acquire subsurface sediments below the approximate elevation of the channel depth of -25 feet MLLW.

At each location, a 15-foot-long vibracore will be advanced until it can penetrate no further. Each core will be logged and sectioned into approximately 1- to 2-foot intervals (no interval will be greater than 2 feet), unless otherwise indicated based on visual observation and stratigraphy, for testing. Because surface sediment quality will have already been characterized (cores are collocated with surface samples), only two subsurface intervals per core will initially be submitted for physical and chemical testing. The core intervals targeted for analysis will be based on field screening, and will include one sample (the 'midpoint core interval') exhibiting significant potential for contamination (if observed) and one sample interval below the potentially contaminated sample interval (the 'lower core interval') where no visible indication of potential contamination is observed. The surface, midpoint, and lower core interval data will be used to quantify the vertical extent of contamination and inform contaminated sediment volume evaluations in the FS. All remaining core intervals will be archived for future analysis, if needed.

If NAPL or another obvious form of contamination is identified during the processing of subsurface cores collected at any perimeter subsurface core locations (Appendix B), an additional core will be collected offset from the initial location. If necessary, the exact placement will be made by the Field Coordinator (identified in Appendix B) and reported to EPA.

The planned subsurface investigation will be completed using vibracore exploration methods. To the extent that the findings of upland and sediment investigations indicate that Site-related contamination is likely to be present in sediment strata that could not be evaluated using these sampling methods, other sampling approaches will be considered. If alternative methods (e.g., use of barge-mounted auger drilling methods) are warranted, the methods and locations for such follow-up investigations would be defined in an addendum to this Work Plan (see Section 6.2).

5.5.2.6 Surface Water Investigation

Surface water samples will be collected from two locations within the ISA and at two background locations for testing of physical properties and chemical analysis (Figures 5-6 through 5-8). These data will be used to inform the HHRA and ERA. To assess potential variability associated with seasons and weather conditions, four quarterly sampling events will be conducted. One of the sampling events will target a rain event, and another will target a relatively dry period. At each location, samples will be collected from 3 feet below the water surface and 3 feet above the mudline. The surface water samples will be submitted for an analysis of Site COPCs, conventional parameters, and alkylated PAHs. Field measurements of dissolved oxygen, pH, salinity, and temperature will be recorded at each sampling depth.

5.6 Contingent Studies

Other studies in addition to those described in Section 5.5 may be necessary to characterize the Site for the RI/FS. However, the need and scope of these studies will

depend on the results of the initial studies. Potential contingent studies are described below.

5.6.1 Upland Investigation

Contingent upland investigation activities may be warranted to fill remaining data gaps after completion of the work described in Section 5.5.1. These contingent investigation tasks may include the following:

- Additional investigation into the nature of NAPL, if free-phase NAPL is identified, by applicable petrophysical testing methods;
- Additional investigation into the extent of NAPL by applicable *in-situ* and/or *ex-situ* characterization techniques;
- Sampling of soil vapor and/or indoor air, if the soil and/or groundwater data indicate a potential risk to existing occupied 42 or future structures;
- Evaluation of the groundwater-to-surface water pathway; and
- Development of hydraulic and/or contaminant fate-and-transport groundwater models.

The scope of and methods for these studies, if needed, will depend on the results of the initial investigations and are, therefore, not provided in this Work Plan. An addendum to this Work Plan would be prepared if additional studies are needed. A brief description of potential contingent activities is provided below.

If NAPL is present at sufficient volumes in any wells, bail-down tests may be used to estimate the transmissivity of DNAPL and LNAPL. Other petrophysical testing methods may also be applicable, depending on the type, quantity, and location of NAPL identified during the RI.

The TarGOST® technology, which uses laser-induced fluorescence to delineate manufactured gas plant tar or creosote NAPL (moderate to heavy concentration of PAHs), could possibly be used to detect and characterize NAPL in fill and shallow native soils in areas where manufactured gas plant tar or creosote has been identified by other investigation methods. However, TarGOST® is specifically intended for use in delineating NAPL-contaminated zones and is appropriate for sites where there is a confirmed presence of manufactured gas plant tar or creosote NAPL. In addition, TarGOST® is conducted using direct-push drilling methods that likely have limited depth penetration capabilities at the Site due to the dense glacial soils. A preliminary understanding of the extent to which NAPL is present in shallow or deeper soils at the Site, and a better understanding of the nature of subsurface soils at the Site is needed to determine whether the use of TarGOST® could be successful at the Site.

Ultraviolet (UV) light photography could be used to characterize NAPL occurrence and extent with low to moderate concentrations of PAH components. The technique uses a digital image of a soil core in an area of known or suspected NAPL to evaluate the nature of the NAPL, such as its pore space saturation and its potential mobility. UV light

⁴² Existing structures within the ISA include two warehouse buildings used for storage of automotive parts and equipment. Ongoing use and occupancy of these structures will be further evaluated during the RI to determine if additional assessment of indoor air is warranted.

photography can also determine the relative impacts within a single core to identify the most heavily impacted zone and identify variation in NAPL impacts between soil lithologies within the core.

After the extent of contamination in soil, groundwater, sediment, and surface water is better defined, ongoing transport of contaminants to surface water/sediment will be evaluated. This may include assessment of the continuity of contamination from the upland to marine areas through groundwater and sediment porewater sampling, including depth profiles of contamination in porewater; delineation of NAPL at the shoreline; and estimates of contaminant flux in groundwater.

Hydraulic and/or contaminant transport groundwater models may be useful tools for conducting the RI and FS. These tools can be used in conjunction with empirical data to further the understanding of contaminant fate and transport and support the engineering evaluations of remedial technologies such as groundwater pumping. However, additional Site information is needed to evaluate the usefulness of these tools and which models might be appropriate.

As discussed in Sections 6.1 and 6.2, any contingent work activities will be proposed based on the data gaps identified in the Phase 1 Data Report. The scope of work and sampling methodology for the contingent upland investigation would be described in detail in an addendum to this Work Plan (Section 6.2), which would be approved by EPA before the completion of any additional work.

5.6.2 Marine Investigation

Contingent sediment investigation activities may be warranted to fill remaining data gaps after completion of the work described in Section 5.5.2. These contingent investigation tasks may include the following:

- Potential step-out surface or subsurface sampling in the sediment areas of the Site, if needed to define the nature and extent of Site-related contamination.
- Supplemental subsurface sediment coring using alternative methods, if needed, to
 evaluate the distribution of Site-related contamination not accessible using
 vibracore methods.
- Sediment bioassay and/or porewater testing, if necessary to confirm the estimated extent of benthic infaunal community impacts for the ERA.
- Testing of Site-related contaminant concentrations in tissues in relevant seafood species or prey species where necessary to support the HHRA or ERA.
- Sediment geochronology testing, if it is determined necessary to support the evaluation of sediment stability and natural recovery processes.

The scope of and methods for these studies, if needed, will depend on the results of the initial investigations and are, therefore, not provided in this Work Plan. An addendum to this Work Plan would be prepared if additional studies are needed (see Section 6.2).

6 Remedial Investigation Tasks

This Section provides a general description of the tasks to be performed to complete the RI in accordance with the AOC, the SOW and EPA RI/FS guidance (EPA 1988a). It also summarizes the various phases of work and how each phase relates to the next phase. A flowchart showing the sequence of the remedial investigation and risk assessment components is provided on Figure 1-2. A general schedule for completion of the work including key decision points is provided in Section 8. Specific details of field investigation methods and sampling approaches, as currently planned, are provided in Appendix A.

6.1 Planned Remedial Investigation Activities

The planned work activities, as described in Section 5.5, will be completed to meet the objectives of the RI/FS in accordance with the requirements of the SOW. The collection of data will address the data needs to assess the current and future potential risk to human health and the environment and allow for the development and screening of remedial action alternatives. The planned work activities, presented herein, are those anticipated to be necessary to meet the RI/FS objectives, which are further specified in the SOW:

- Investigate and define the physical, chemical, and biological characteristics of the Site;
- Define the sources of contamination;
- Define the human and ecological uses of Site; and
- Describe the nature and extent of contamination.

Field and preliminary laboratory data will be provided to EPA as it is received, following the field communication plans in Appendices A and B, to enable adaptive management practices in evaluating whether the RI/FS objectives have been met. Data may also be summarized in tabular or visual form as needed to support work planning. Some specifics of the field investigation, such as the number, location, and depth of step-out trenches, borings and wells, will be determined during the investigation following the decision-making criteria outlined in this Work Plan. Proposed and completed field activities will also be communicated in accordance with the field communication plans to enable ongoing review and input from the Cascade and EPA project teams.

After the completion of the work activities described in this Work Plan, the Phase 1 Data Report will be prepared to compile the collected data. In accordance with the SOW, the Phase 1 Data Report will describe and display information and data collected during the Site characterization activities, including the sampling locations and the distribution of contaminant concentrations. If data needs are identified that require activities not covered by this Work Plan, one or more addenda to this Work Plan may be prepared (see Section 6.2).

6.2 Contingent Remedial Investigation Activities

If determined to be necessary to satisfy outstanding data needs and meet the objectives of the RI/FS, contingent studies will be proposed in one or more addenda to this Work Plan. The contingent studies may consist of the expansion of previous studies, potential

contingent studies identified in Section 5.6, or other studies that are warranted based on the collected data. Addenda to this Work Plan, if applicable, may be submitted with the Phase 1 Data Report. Each addendum to this Work Plan will present the proposed scope of work, including the basis for the additional work and the rationale for the sampling locations and/or methodology.

Data collected during contingent studies will be documented and submitted to EPA in the Phase 2 Data Report.

6.3 Risk Assessment

The RI/FS will include collection of information and data necessary to perform a baseline HHRA and ERA, in accordance with the SOW (EPA, 2013a). The risk assessment strategy will be developed in consultation with EPA and the Tribe. The risk assessment will consider current and potential future land uses at the Site, considering local land use designations applicable to the Former Gas Works Property and the Sesko Property. A Risk Assessment Technical Memorandum will be prepared in conjunction with the Phase 1 Data Report to present the preliminary screening of the RI data and provide a detailed description of the methods to be used for the baseline risk assessments. The Risk Assessment Technical Memorandum will evaluate the data presented in the Phase 1 Data Report. The Risk Assessment Technical Memorandum will be submitted to EPA for concurrence that sufficient data has been collected, or to propose the collection of additional data, to enable preparation of the draft baseline HHRA and ERA. The scope and key elements of the HHRA and ERA are described in Section 5.3.

The draft reports for the baseline HHRA and ERA will be submitted to EPA as part of the Draft RI Report (Section 6.4). After EPA has reviewed the Draft RI Report and provided comments, the final risk assessment reports will be submitted to EPA with the Final RI Report (Section 6.4).

6.4 Remedial Investigation Report

After the completion of any contingent studies and EPA approval of the data report summarizing the final phase of investigation (either the Phase 1 Data Report or a Phase 2 Data Report), a Draft RI Report will be prepared to summarize the results of all phases of the field activities conducted to characterize the contaminant sources, evaluate the nature and extent of contamination, and evaluate the fate and transport of contaminants. The Draft RI Report will be submitted to EPA for review in accordance with the requirements of the AOC. After the receipt of EPA comments, a Final RI Report will be prepared.

7 Feasibility Study Planning

Section 7 describes tasks involved in developing the FS, particularly the review of remedial approaches, including potentially applicable remedial technologies, based on similar sites. Further development of the CSM is necessary before identifying remedial approaches for the Site. A preliminary review of the remedial technologies helps determine what data may need to be collected during the field investigations discussed in Section 5.

7.1 Feasibility Study Tasks

This Section describes the tasks that will be conducted to identify potential remedial approaches and select a remedy for the Site in accordance with the AOC, the SOW, and EPA RI/FS guidance (EPA 1988a). This Section also summarizes each phase of the FS and how each phase relates to the next one. A tentative schedule for completion of the FS is provided in Section 8.

7.1.1 Remedial Alternatives Development/Screening

The first step in the FS process will be the preparation of an Alternatives Development Memorandum that identifies and screens a range of potential remedial alternatives to determine whether they should be included in a more detailed analysis. The Alternatives Development Memorandum will include the following:

- Identification of refined RAOs based on the results of the RI and baseline risk assessments:
- Development of general, potential response actions for each medium of interest to meet the RAOs;
- Identification of areas and volumes of Site-related COPCs to which the general response actions may apply;
- Identification and evaluation of remedial technologies applicable to each general response action and a screening to determine and document those that will be eliminated from further evaluation;
- A presentation of the selected remedial technologies and their assembly into remedial action alternatives for the Site:
- A summary of the action-specific and contaminant-specific ARARs and PRGs for each of the assembled remedial action alternatives;
- A screening of the assembled remedial action alternatives based on short- and longterm effectiveness, implementability, and relative cost, if necessary.

The Alternatives Development Memorandum will be prepared after EPA approval of the Final RI Report.

7.1.2 Treatability Study/Pilot Testing

Treatability studies and/or pilot testing of potential remedial technologies will be performed after the preparation of the Alternatives Development Memorandum, if necessary to support further evaluation of the retained alternatives. If treatability studies or pilot testing are determined to be necessary to evaluate a technology, a Treatability Testing Work Plan will be prepared to describe the technology, present the purpose of the treatability study/pilot testing, and summarize the testing approach and methodology, including a Sampling and Analysis Plan, if appropriate. The results of the treatability study/pilot testing will be summarized in a Treatability Study Evaluation Report, which will be submitted to EPA as a draft for review and comment; any comments provided by EPA will be addressed in a final version of the report.

7.1.3 Detailed Analysis of Alternatives

A detailed analysis of the final set of alternatives (Section 7.1.1) and the results of any treatability studies and/or pilot testing (Section 7.1.2) will be performed. It will consist of an analysis of each alternative in terms of nine CERCLA evaluation criteria (EPA 1998a) and a comparative analysis of all the alternatives using the same criteria as a basis for comparison. The results will be documented in an Alternatives Evaluation Memorandum.

7.1.4 Feasibility Study Report

After the receipt of EPA comments on the Alternatives Evaluation Memorandum, the Draft FS Report will be prepared to present the basis for remedy selection and document the development and analysis of the remedial alternatives. The Draft FS Report will be submitted to EPA for review in accordance with the requirements of the AOC. After the receipt of EPA comments, a Final FS Report will be prepared.

7.2 Potential Remedial Approaches

This Section describes potentially applicable remedial technologies and approaches based on similar sites. The purpose of this description is to provide an initial understanding of what remedial technologies may be applied and identify data required to evaluate the feasibility of each technology to meet the RAOs. The selected remedy is typically a combination of multiple remedial technologies to achieve all RAOs.

7.2.1 Potential Remedial Technologies

Achieving RAOs at a site typically occurs by implementing a combination of several remedial technologies. Depending on site-specific circumstances, the selected remedial technologies may result in the complete elimination or destruction of hazardous substances at the site, the reduction or elimination of migrating hazardous substances at the site, or some combination of these effects. Remedial technologies may be used in combination with engineering controls (e.g., barriers such as fences or caps) or institutional controls (i.e., non-engineered controls such as land use restrictions) when hazardous wastes remain at a site. The National Contingency Plan states a preference for remedial technologies at Superfund sites that involve treatment, including but not limited to controlling or eliminating sources and reducing or eliminating exposure pathways, particularly for highly contaminated materials.

Remedial technologies are often categorized by the following general response actions, which are applicable to both upland and marine environments:

- Monitored Natural Attenuation. Natural attenuation is the reduction of contaminant concentrations at the point of exposure over time by means of natural processes, such as sedimentation, sorption, dispersion, and/or biodegradation. Monitoring documents that the processes are occurring at the desired rates. For sediment, this general response action is referred to as monitored natural recovery.
- *In Situ* Containment. *In situ* containment involves confining hazardous substances in place by the placement of physical barriers or hydraulic controls. Containment technologies can be designed to prevent contact with and/or migration of hazardous substances.
- *In Situ* Treatment. *In situ* treatment technologies can potentially reduce the concentration, mobility, and/or toxicity of COCs.
- **Removal.** Contaminated materials can be physically removed from a site and treated and/or disposed of at either an on-site or an off-site permitted disposal facility.
- Ex Situ Treatment. Ex situ treatment technologies destroy or immobilize contaminants in media that have been removed from the subsurface.
- **Disposal.** Disposal technologies include the placement of contaminated solid media in on-site or off-site landfills or the discharge of contaminated water to a publicly owned treatment works.

Preliminary lists of potential remedial technologies for NAPL, soil, groundwater, and sediment at the Site are provided in Tables 7-1 through 7-4, respectively.

7.2.2 Remedial Approaches at Other MGP Sites

Hundreds of MGP sites around the country have been through or are undergoing an RI/FS and cleanup action. Table 4-2 identifies remedial approaches that have been fully or partially implemented at MGP sites with characteristics (e.g., geology and presence of adjacent surface water bodies) that are similar to the Bremerton Gas Works Superfund Site. Common actions have included combinations of removal with off-site disposal or on-site treatment, solidification/stabilization, and institutional and engineering controls. Other technologies have included pump-and-treat, bioremediation, *in situ* chemical oxidation, barriers, and NAPL collection.

7.2.3 Feasibility Study Data Gaps

As part of the FS, the potential remedial technologies identified in Tables 7-1 through 7-4 will be evaluated based on each technology's effectiveness, ability to be implemented, cost and fit. Evaluating and analyzing each into remedial technology requires a good understanding of Site characteristics. In general, data gathered during the RI to develop the CSM (including characterizing physical characteristics of the Site, e.g., hydrogeology and groundwater flow, delineating the nature and extent of contamination, evaluating contaminant fate and transport, and assessing risks to human health and the environment) will also support the development and evaluation of remedial alternatives.

Data from characterization of the Site will need to be sufficient to develop hydraulic and contaminant fate-and-transport models that may be needed to assist in the engineering evaluations during the FS (e.g., in developing and evaluating alternatives that use groundwater extraction or dewatering). The data will also need to be sufficient to delineate the extent of contaminant source areas or "hot spots."

In addition, the following information is anticipated to be necessary to complete the FS:

- Geotechnical data (e.g., for developing excavation and shoring plans), including penetration test data, soil moisture content, Atterberg limits, and gradation;
- Recoverability characteristics of NAPLs, if present;
- Waste characteristics (e.g., toxicity characteristic leaching procedure [TCLP]) to determine potential disposal and/or treatment options; and
- Evaluations of current velocity and sediment substrate study by means of a towed video camera, to evaluate physical forces and geologic formations.

Additional technology-specific data needs may be identified as more data are collected and the FS alternatives are developed. These may include Site characterization data, bench testing, or pilot testing of potential remedial technologies.

8 Schedule

The field investigation activities described herein will commence within 30 days after receipt of EPA's written approval of this Work Plan. The estimated schedule and sequencing of field investigation activities is provided in Table 8-1. The actual schedule may vary based on several factors including contractor availability, the date EPA approves this Work Plan, and adjustments to the scope of work based on field investigation findings.

The schedule for completing RI/FS investigation activities and deliverables will be consistent with the deadlines defined in the AOC, which include the following:

- Prepare and submit the Phase 1 Data Report to EPA within 90 days after completion
 of Site characterization activities and receipt of final validated data. The Phase 1 Data
 Report will summarize the results of the Site characterization activities and identify
 any outstanding data needs.
- If warranted by the results summarized in the Phase 1 Data Report, prepare an addendum to this Work Plan describing the additional Site characterization activities necessary to meet the objectives of the RI/FS. After EPA approval of the addendum to this Work Plan, complete the additional Site characterization activities.
- Prepare and submit a Phase 2 Data Report to EPA within 90 days after completion of the additional Site characterization activities and receipt of final validated data. The Phase 2 Data Report will summarize the results of the additional Site characterization activities.
- The Draft Baseline Ecological and Human Health Risk Assessment Reports will be prepared and submitted to EPA within 180 days after receipt of all final validated data obtained during Site characterization activities, including any contingent studies.
- The Draft RI Report will be prepared and submitted to EPA within 360 days after receipt of all final validated data obtained during Site characterization activities, including any contingent studies.
- The Final RI Report, which will include the Final Baseline Ecological and Human Health Risk Assessment Reports, will be submitted to EPA within 90 days after receipt of comments from EPA on the Draft RI Report.
- The Alternatives Development Memorandum will be submitted to EPA within 90 days after receipt of EPA's written approval of the Final RI Report.
- If necessary, a Treatability Testing Work Plan, treatability testing, and the Treatability Study Evaluation Report will be completed to further evaluate alternatives introduced in the Alternatives Development Memorandum. A separate schedule will be prepared for these activities if they are deemed necessary.
- The Alternatives Evaluation Memorandum will be submitted to EPA within 90 days after receipt of EPA's comments on the Alternatives Development Memorandum and Treatability Study Evaluation Report, if applicable.
- The Draft FS Report will be submitted to EPA within 120 days after receipt of EPA's written approval on the Alternatives Evaluation Memorandum.



9 Project Management Plan

This Section identifies key project staff and responsibilities and describes lines of communication and project coordination details. It also includes a description of data management procedures.

9.1 Project Management

The RI/FS is being conducted by Cascade. EPA is providing regulatory oversight of the RI/FS activities in accordance with the AOC. Cascade and EPA project teams are described in Section 9.1.1.

Other participating entities (i.e., stakeholders) include property owners, other regulatory agencies, and interested organizations. Key stakeholders who participate in the scoping, review, and comment on the RI/FS, and their general roles, are described in Section 9.1.2. EPA has the primary responsibility for engaging and coordinating with these key stakeholders throughout the RI/FS process and Cascade and its contractors will provide support, as needed, for these coordination efforts.

9.1.1 Project Teams

The Remedial Project Manager (RPM) for EPA is:

Eva DeMaria
EPA Region 10, Office of Environmental Cleanup (ECL-113)
1200 Sixth Avenue

Seattle, Washington 98101 Phone: (206) 553-1970

E-mail: DeMaria.Eva@epa.gov

The Project Coordinator for Cascade is:

Kalle Godel Montana-Dakota Utilities Co. 400 North Fourth Street Bismarck, North Dakota 58501

Phone: (701) 222-7657

E-mail: Kalle.Godel@mdu.com

The Cascade Project Coordinator is responsible for administering the actions required by the AOC.

EPA's oversight contractor for the RI/FS work is CH2M. The project manager for CH2M is Susan Moore.

Cascade's consultant project team consists of representatives from Aspect and AnchorQEA and their subconsultants and subcontractors. Aspect will be coordinate RI/FS activities for the upland area of the Site. Anchor QEA will coordinate RI/FS activities in the marine area of the Site and conduct the risk assessment. Aspect will be responsible for overall project management and production of RI/FS deliverables.

The project managers for Aspect and Anchor QEA, who have final authority and responsibility for their teams' activities, are as follows:

• **Aspect:** Jeremy Porter

Anchor QEA: Mark Larsen

Supporting project team members and team management structure for conducting the Site characterization activities described in this Work Plan are provided in the Upland and Marine SQAPPs (Appendices A and B).

All work will be conducted in accordance with the consultants' Quality Management Plans, which have been previously submitted to EPA in accordance with Section VIII of the AOC.

All work conducted by Aspect and Anchor QEA will be completed in accordance with applicable state and federal worker health and safety requirements. The site-specific Health and Safety Plans for each organization, which establishes the procedures and practices to protect their workers from potential hazards posed by field activities at the Site, are included as Appendices G (Aspect) and H (Anchor QEA).

9.1.2 Stakeholders

Key stakeholders and their general roles are as follows:

- The Suquamish Tribe: As described in Section 2.2, the Site and vicinity are in the traditional territory of the Tribe. Under the 1855 Treaty of Point Elliott, the Tribe retained "the right of taking fish at usual and accustomed grounds and stations," which include the Port Washington Narrows. EPA will consult with the Tribe throughout the RI/FS regarding issues of potential interest to the Tribe including matters affecting fish and wildlife or potential cultural or archaeological resources.
- Owners of Property Within the ISA:
 - McConkey Family Trust (owner of the McConkey Property);
 - o Penn Plaza Storage LLC (owner of the Penn Plaza Property);
 - Natasha Sesko (owner of the Sesko Property);
 - o DNR (owner of aquatic tidelands); and
 - o City of Bremerton (owner of public rights-of-way).
- Regulatory and Resource Agencies with jurisdiction at the Site:
 - Kitsap Department of Health (KDOH): A local agency that manages numerous local regulations and programs targeted at human health protection. KDOH has been working with other agencies and stakeholders to improve water and shellfish quality within Port Washington Narrows and Dyes Inlet.
 - Ecology: A state agency responsible for implementing numerous state regulations addressing soil, groundwater and surface water quality,

including but not limited to the Model Toxics Control Act (MTCA; WAC Chapter 173-340), the Sediment Management Standards (SMS; WAC Chapter 173-204) and Washington's Water Quality Standards (WAC 173-201a).

- WDFW: A state agency responsible for implementing Washington State regulations and programs related to protection, enhancement and harvesting of wildlife, fish and shellfish resources.
- WDOH: A state agency responsible for regulations and programs targeted at human health protection, including but not limited to the management of fish and shellfish consumption advisories. The WDOH has also conducted a preliminary health screening of the Site (WDOH 2014).
- NOAA and the United States Fish and Wildlife Service: Federal agencies responsible for implementing Federal regulations and programs related to protection, enhancement and harvesting of wildlife, fish and shellfish resources.

Stakeholders will be provided with periodic communications and invited to meetings regarding the Site throughout the RI/FS process (see Section 9.2), Key draft AOC deliverables, as determined by EPA, will be provided for stakeholder review and opportunity for comment.

Site access for field activities will be coordinated with property owners.

9.1.3 Public Involvement

The public will be engaged and kept informed of Site activities in accordance with the Community Involvement Plan (EPA, Date) developed for the Site. EPA will serve as the lead for public engagement efforts, with the support of Cascade and its contractors.

The Community Involvement Coordinator for EPA is:

Debra Sherbina EPA Region 10, Office of Environmental Cleanup (ECL-113) 1200 Sixth Avenue Seattle, Washington 98101 Phone: (206) 553-0247

E-mail: sherbina.debra@epa.gov

9.2 Project Communications

Periodic communications between the RPM, the Project Coordinator, and the consultants are conducted to minimize delays and to facilitate identification and resolution of potential problems. Project communications include:

• **Progress Reports.** In accordance with the AOC, quarterly progress reports are due to EPA by the 15th day of the month following each quarter. The current schedule involves submittal of progress reports by January 15th, April 15th, July 15th, and October 15th of each year.

- Meetings and Teleconferences. In accordance with the AOC, monthly status calls or meetings are conducted with EPA, unless EPA and Cascade agree to cancel or postpone. Additional meetings and teleconferences are conducted on an as-needed basis. Additional meetings or teleconferences may be held with EPA in presenting initial findings of the RI/FS and risk assessment, evaluating data evaluation approaches, assessing data gap fulfillment, and reviewing deliverables.
- **Stakeholder Briefings.** In accordance with the AOC, periodic briefings on the work will be coordinated with EPA and project stakeholders.
- **Notifications.** In accordance with the AOC, Cascade will notify EPA a minimum of two weeks prior to planned field activities.
- Field Activities and Data Communication. As described in the field communication plan of the Upland and Marine SQAPPs (Appendices A and B), field activities and data (including daily logs, photos, maps/sketches, and monitoring forms) and preliminary analytical data (including pre-validated laboratory reports and, if needed for decision making, summary data tables and/or figures) will be posted to a password-controlled website for review by Cascade and EPA project teams. The website will be set up, tested, and approved by EPA and Cascade prior to beginning field work. During periods of non-routine 43 field activities, a weekly status call will be scheduled with EPA and Cascade project teams. Additional meetings or teleconferences to discuss field activities and preliminary data may be scheduled as needed.

9.3 Data Management

Considerable quantities of data have already been obtained and will be collected during the RI field investigation. This data will need to be stored, checked for quality, and presented in reports. This Section outlines how these data will be managed.

Software and procedures are in place to effectively and efficiently handle data generated during the RI. These systems and processes will ensure that data (e.g., sample numbers, methods, qualifications, locations, etc.) are readily accessible and accurately maintained. The primary steps/elements in the data management process are:

- EarthSoft EQuIS 6 environmental chemistry database setup
- gINT geological boring log database setup
- Sample and analysis planning
- Sample collection

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⁴³ Routine activities include periodic site inspections and groundwater monitoring events.

- Field measurements
- Documentation of location of field activities (GPS, survey, etc.)
- Laboratory analytical data management
- Preliminary reporting and data QA/QC
- Formal data validation (details provided in the SQAPPs) and associated database updates
- Development of maps and tables from EQuIS database, integrated with GIS software as appropriate, to support RI/FS reporting requirements
- Analytical data submittals in accordance with USEPA's Region 10 Data Submission Process for WQX Compatible Deliverables
- Geographic Information Systems (GIS) data submittals in accordance with U.S. EPA Region 10 GIS Data Deliverable Guidance (ed. March 2013)

Data will be collected and recorded in a variety of ways during this project. These include standard field forms (e.g., field data sheets, chain-of-custody forms, and boring logs) and laboratory-generated analytical data. Information about exploration locations, samples, laboratory tests, field measurements and analytical results will be maintained in an EarthSoft EQuIS 6 database. These data will be loaded to EQuIS from electronic data deliverables (EDDs) and preliminarily checked for completes and fidelity against associated reports and documentation. Lithological data will be entered into the gINT database from boring logs under supervision by professional geologists. Access to the EQuIS and gINT databases will be limited to trained project personnel, and the ability add or change data will be granted to only those trained, professional data managers, chemists, and geologists.

Lab reports and other source documents (including original laboratory EDDs) will be filed electronically according to the project-specific storage and retention policies. All electronic data (including the EQuIS and gINT databases) will be backed up nightly in accordance with industry practices.

Data validation will be performed in accordance with the project SQAPPs. Data validation reports will be filed electronically (along with other source documents) and any associated updates to analytical data (including qualifiers and other validation notes) will be added/updated in EQuIS, as appropriate.

10 References

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TABLES

Table 2-1 - Monitoring Well Construction Information and Groundwater Elevation Measurements

Bremerton Gas Works Superfund Site Bremerton, Washington

| | | | | | | | Depth to Water (feet below TOC) | | |
|------------------------|--------------|-------------------|--------------------------------------|------------------------------|----------------------------------|-------------------------------------|------------------------------------|----------|--|
| Well Identification | Installed By | Date Installed | Surface Elevation (Datum Unknown) | Total Boring Depth (Feet) | Depth to Top of Screen (Feet) | Depth to Bottom of Screen (Feet) | 1-Jun-07 | 1-Jun-07 | |
| MP-04 | E&E | 5/13/2008 | 12.38 | 40 | 30 | 40 | | | |
| SP-02 | E&E | 5/12/2008 | 10.44 | 35 | 25 | 35 | | | |
| | | | Surface Elevation in feet (NAVD 88) | | | | | | |
| MW-1 | GeoEngineers | 5/21/2007 | 45.03 | 46.5 | 30 | 45 | 34.68 | 10.35 | |
| MW-2 | GeoEngineers | 5/21/2007 | 42.54 | 46.5 | 30 | 45 | 35.25 | 7.29 | |
| MW-3 | GeoEngineers | 5/22/2007 | 39.1 | 46.5 | 30 | 45 | 32.9 | 6.2 | |
| MW-4 | GeoEngineers | 5/23/2007 | 35.2 | 41.5 | 20 | 40 | 29.32 | 5.88 | |
| MW-5 | GeoEngineers | 5/24/2007 | 18.51 | 21.5 | 5 | 20 | 15.21 | 3.3 | |
| MW-6 | GeoEngineers | 5/22/2007 | 34.95 | 36.5 | 15 | 35 | 30.2 | 4.75 | |
| MW-7 | GeoEngineers | 5/23/2007 | 33.24 | 36.5 | 15 | 35 | 30.21 | 3.03 | |
| MW-8 | GeoEngineers | 5/22/2007 | 35.56 | 41.5 | 20 | 40 | 32.64 | 2.92 | |

Notes:

-- = not measured E&E = Ecology and Environment NAVD 88 = North American Veritcal Datum of 1988 TOC = top of casing

Table 3-1 – Potential ARARs, Chemical-Specific

Bremerton Gas Works Superfund Site Bremerton, Washington

| Act/Authority | Criteria/Issue | Citation | Brief Description | Applicability/Appropriateness |
|---|---|--|---|--|
| Safe Drinking Water Act | Federal Primary Drinking Water Standards – MCLs and MCLGs | 42 USC 300f; 40 CFR 141, Subpart O | Establishes drinking water standards for public water systems to protect human health. Includes standards for the following Site chemicals of concern: arsenic, benzene, and benzo(a)pyrene. The National Contingency Plan states that MCLs, not MCLGs, are ARARs for usable aquifers. | ARARs for groundwater that could potentially be used for drinking water, where the water will be provided directly to 25 or more people or will be supplied to 15 or more service connections. |
| Safe Drinking Water Act | Federal Secondary Drinking Water Standards – Secondary MCLs | 42 USC 300f; 40 CFR 143 | Establishes drinking water standards for public water systems to achieve the aesthetic qualities of drinking water (secondary MCLs). | TBC for groundwater that could potentially be a drinking water source (i.e., achieved as practicable). |
| Clean Water Act | Federal Ambient Water Quality Criteria | 33 USC 1311– 1317; 40 CFR 131 | Under Clean Water Act, Section 304(a), minimum criteria are developed for water quality programs established by states. Two kinds of water quality criteria are developed: one for protection of human health, and one for protection of aquatic life. The federal recommended water quality criteria are published on EPA's website: http://water.epa.gov/scitech/swguidance/standards/current/index.cfm | ARARs for surface water if more stringent than promulgated state criteria. |
| EPA Superfund Soil Screening Guidance | Risk-Based, Site-Specific Soil Screening Levels (SSLs) | | Provides guidance with a tiered framework for developing risk-based, site-specific soil screening levels (SSLs) for the protection of human health. | ARARs for soil |
| Surface Water Quality Standards | State Ambient Water Quality Criteria | Chapter 90.48 RCW; Chapter 173-201A WAC | Establishes water quality standards for protection of human health and for protection of aquatic life (for both acute and chronic exposure durations). | ARARs for surface water where Washington State has adopted, and EPA has approved, water quality standards. |

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Table 3-1 – Potential ARARs, Chemical-Specific

Bremerton Gas Works Superfund Site Bremerton, Washington

| Act/Authority | Criteria/Issue | Citation | Brief Description | Applicability/Appropriateness |
|-------------------------------------|---|---|--|--|
| Model Toxics Control Act | State Soil, Air, Groundwater, and Surface Water Cleanup Standards | Chapter 70.105D RCW; Chapter 173-340 WAC | Establishes cleanup levels for Site groundwater, surface water, soil, and air, including rules for evaluating cross-media protectiveness for all potential receptors (humans and terrestrial plants and animals). MTCA cleanup levels cannot be set at concentrations below natural background. | Promulgated numeric cleanup levels are ARARs for soil, air, groundwater, and surface water. Equations to develop cleanup levels are not ARARs. |
| Sediment Management Standards | State Sediment Quality Criteria | Chapters 90.48 & 70.105D RCW; Chapter 173-204 WAC | Establishes both numerical and biological wasting-based standards for the protection of benthic invertebrates in marine sediments. The current rule also defines methods for establishing cleanup levels protective of human health, including protection from risks associated with seafood consumption, analytical considerations, and natural and regional background contamination levels. | SMS cleanup levels will serve as ARARs for the development of sediment cleanup levels. |

Notes:

ARAR = applicable or relevant and appropriate requirement

CFR = Code of Federal Regulations

EPA = U.S. Environmental Protection Agency

MCL = maximum contaminant level

MCLG = maximum contaminant level goal

MTCA = Model Toxics Control Act

RCW = Revised Code of Washington

SMS = Sediment Management Standards

TBC = to be considered

USC = United States Code

WAC = Washington Administrative Code

Table 3-2 – Potential ARARs, Location-Specific

Bremerton Gas Works Superfund Site Bremerton, Washington

| Act/Authority | Criteria/Issue | Citation | Brief Description | Applicability/Appropriateness |
|---|---|---|--|--|
| Endangered Species Act | Effects on Endangered Species | 16 USC 1531 et seq.; 50 CFR 17 | Actions authorized, funded, or carried out by federal agencies may not jeopardize the continued existence of endangered or threatened species or adversely modify or destroy their critical habitats, or must take appropriate mitigation steps. | ARAR for remedial actions that may adversely affect endangered or threatened species or critical habitat present at the Site. |
| Treaty of Point Elliott, 1855 | Tribal lands, hunting, and fishing rights | Articles of agreement and convention made and concluded at Point Elliott, in the territory of Washington. Ratified March 8, 1859. Proclaimed April 11, 1859. | Article 5 secures Tribal right of taking fish at usual and accustomed grounds and stations. | ARAR for fish for shellfish harvest in and adjacent to the Site. |
| Safe Drinking Water Act | Underground Injection Control, Sole Source Aquifer Program, and Wellhead Protection Program | 42 USC 300h–300h-8; 40 CFR 300.400(g)(4); Chapter 173-160 WAC; WAC 246-290- 135 | Resource planning programs designed to prevent contamination of underground sources of drinking water. | The requirements of the City's wellhead protection program are TBCs as a performance standard for groundwater that is a potential drinking water source (i.e., achieved as practicable). (Note that there are no water supply wells near the Site that are currently regulated by the City's program.) |
| Migratory Bird Treaty Act of 1918 | Protection of Migratory Birds | 16 USC 703-712 50 CFR 10.13 | Makes it illegal to take, possess, sell, purchase or barter any migratory bird except under the terms of a valid permit. | ARAR if migratory birds are impacted during investigation or remedial action. |

Table 3-2 – Potential ARARs, Location-Specific

Bremerton Gas Works Superfund Site Bremerton, Washington

| Act/Authority | Criteria/Issue | Citation | Brief Description | Applicability/Appropriateness |
|--|---------------------------------|--|--|--|
| Magnuson- Stevens Fishery Conservation and Management Act | Habitat Impacts | 16 USC 1855(b); 50 CFR 600.920 | Requires evaluation of impacts on EFH if activities may adversely affect EFH. | ARAR if the remedial action may adversely affect EFH. |
| Executive Order for Wetlands Protection | Wetlands Impacts | Executive Order 11990 (1977), 40 CFR 6.302(a); 40 CFR 6, App. A | Requires measures to avoid adversely affecting wetlands whenever possible, to minimize wetland destruction, and to preserve the value of wetlands. | ARAR for assessing impacts on wetlands, if any, from the remedial action and for developing appropriate compensatory mitigation. |
| Marine Mammal Protection Act, and the | Protection of Marine Mammals | 16 USC Chapter 31 | Prohibits the taking (to hunt harass, capture, or kill) of marine mammals in U.S. waters. | ARAR if marine mammals are impacted during investigation or remedial action. |
| Marine Protection, Research, and Sanctuaries Act | Ocean Dumping | 16 USC § 1431 et seq. and 33 USC §1401 et seq. | Prohibits transportation of material from the U.S. for purpose of ocean dumping; transportation of material from anywhere for the purpose of ocean dumping by U.S. agencies or U.Sflagged vessels; dumping of material transported from outside the U.S. into the U.S. territorial sea except under the terms of a valid permit. | ARAR if ocean dumping is planned to occur, except under the terms of a valid permit. |

Table 3-2 – Potential ARARs, Location-Specific

Bremerton Gas Works Superfund Site Bremerton, Washington

| Act/Authority | Criteria/Issue | Citation | Brief Description | Applicability/Appropriateness |
|--|---|-----------------------------|---|--|
| Fish and Wildlife Conservation Act ("Nongame Act") | Conservation of nongame fish and wildlife | 16 USC 2901-2911 | Authorizes financial and technical assistance to the States for the development, revision, and implementation of conservation plans and programs for nongame fish and wildlife. | ARAR if conservation plans and programs for nongame fish and wildlife are implemented. |
| Bald Eagle Protection Act | Protection of Bald and Golden Eagles | 16 USC 668(a); 50 CFR 22 | Prohibits the take, possession, sale, purchase, barter, transport or import, of any bald or golden eagle, alive or dead, including any part, nest, or egg, except under the terms of a valid permit | ARAR if bald or golden eagles are impacted during investigation or remedial action. |

Notes:

ARAR = applicable or relevant and appropriate requirement

CFR = Code of Federal Regulations

City = City of Bremerton

EFH = essential fish habitat

TBC = to be considered

USC = United States Code

WAC = Washington Administrative Code

Table 3-3 – Potential ARARs, Action-SpecificBremerton Gas Works Superfund Site

| Remedial Activity | Act/Authority | Criteria/Issue | Citation | Brief Description | Applicability/Appropriateness |
|---------------------|--|--|---|--|---|
| Soil Excavation and | Solid Waste Disposal Act | Management and Disposal of Solid Waste | 42 USC 6901–6917; 40 CFR 257–258 | Establishes requirements for the management and disposal of solid wastes. | ARAR for remedial actions that result in upland disposal of excavated or dredged material. |
| Upland Filling | Resource Conservation and Recovery Act (RCRA); Washington Hazardous Waste Management Act and Dangerous Waste Regulations | Generation and Management (Transportation, Treatment, Storage, and Disposal) of Hazardous Waste; Off-Site Land Disposal Considerations | 42 USC 6921–22; 40 CFR 260, 261, and 268; Chapter 70.105 RCW; Chapter 173-303 WAC (Chapter 173-307 WAC Pollution Prevention Plans is a TBC) | Defines solid wastes subject to regulation as hazardous wastes. Requires management of hazardous waste from "cradle to grave" unless exemption applies. MGP wastes are subject to certain exemptions (e.g., Bevill Amendment provisions) | ARAR for wastes and soils sediments excavated from the Site for off-site disposal, and a TBC for on-site stabilization or containment actions. |
| | Hazardous Materials Transportation Act | Transport of Hazardous Materials | 49 USC 5101 et seq.; 49 CFR 171–177 | Establishes requirements for transport of hazardous materials. | ARAR for those hazardous materials (e.g., DNAPL) transported off site. |
| | Washington Hydraulics Code | Filling of Wetlands | Chapters 75.20 and 77.55 RCW; Chapter 220-110 WAC | Establishes requirements for performing work that would alter existing jurisdictional wetlands. | ARAR if remedial actions such as excavation or capping affect existing jurisdictional wetlands. Remedial actions must result in no net loss of aquatic habitat and function after sequential consideration of avoidance and mitigation, allowing for site-specific evaluations of existing wetland functions. |

Table 3-3 – Potential ARARs, Action-Specific

Bremerton Gas Works Superfund Site

| Remedial Activity | Act/Authority | Criteria/Issue | Citation | Brief Description | Applicability/Appropriateness |
|--|--|---|--|---|---|
| Soil Excavation and Upland Filling (Continued) | City of Bremerton Shoreline Master Program and Critical Areas Regulations | Shoreline of Statewide Significance; Fish and Wildlife Habitat Conservation Areas | Chapter 90.58 RCW; Chapter 173-14 WAC; City of Bremerton Ordinance #5299 (effective December 4, 2013); Critical Area Regulations (BMC 20.14) are incorporated into the SMP by reference | Establishes replacement requirements for FWHCAs affected by remedial actions to ensure no net loss of existing ecological function; also establishes requirements for buffers and setbacks from shorelines. | ARAR if remedial actions such as excavation or capping result in impacts within 200 feet of ordinary high water mark or designated FWHCAs. Remedial actions must result in no net loss of aquatic habitat and function after sequential consideration of avoidance and mitigation, allowing for site-specific evaluations of existing shoreline habitat and FWHCAs. Washington's vested rights rule governs which SMP requirements apply in a given circumstance. Substantive requirements of the SMP that were in effect when redevelopment project applications were filed may be ARARs for future redevelopment actions at the Site. |
| Dredging, Capping, and/or Discharge to Puget Sound | Clean Water Act | Federal Ambient Water Quality Criteria | 33 USC 1311–1317; 40 CFR 131 | Regulates activities that may result in discharges into navigable waters. | ARAR for control of short-term impacts on surface water due to implementation of remedial actions that include dredging, capping, and discharge of treated water into Puget Sound. Incorporates the substantive provisions of relevant and appropriate Joint Aquatic Resources Permit Application (JARPA), Nationwide Permit, and stormwater regulation requirements. |

Table 3-3 – Potential ARARs, Action-Specific

Bremerton Gas Works Superfund Site

Bremerton, Washington

| Remedial Activity | Act/Authority | Criteria/Issue | Citation | Brief Description | Applicability/Appropriateness |
|---|--|---|---|--|---|
| | Surface Water Quality Standards | State Ambient Water Quality Criteria | Chapter 90.48 RCW; Chapter 173-201A WAC | Regulates activities that may result in discharges into navigable waters. | ARAR for control of short-term impacts on surface water sue to implementation of remedial actions that include dredging, capping, and discharge of treated water into Puget Sound. Incorporates the substantive provisions of relevant and appropriate requirements, where Washington State has adopted, and EPA has approved, water quality standards. |
| Dredging, Capping, | Clean Water Act | Discharge of Materials into Puget Sound | 33 USC 1344; 40 CFR 230 | Regulates discharge of dredged and fill material into navigable waters of the United States. | ARAR for dredging and capping activities in Puget Sound. |
| and/or Discharge to Puget Sound (Continued) | Clean Water Act | Discharge of Materials into Puget Sound | 33 USC 1251; 40 CFR 122, 123 and 124 | Implements the National Pollutant Discharge Elimination System (NPDES) Program, which regulates discharge of pollutants from any point source into waters of the United States. | ARAR for remedial actions if construction stormwater and/or treated water is discharged into Puget Sound. |
| | Fish and Wildlife Coordination Act Discharge of Materials, Impoundment or Diversion of Waters in Puget Sound | | 16 USC 662 and 663; 40 CFR 6.302(g) | Requires federal agencies to consider effects on fish and wildlife from projects that may alter a body of water and mitigate or compensate for project-related losses, which include discharges of pollutants to water bodies. | ARAR for in-water remedial actions or if treated water is discharged into Puget Sound. |
| | River and Harbors Act Placement of Structures in Puget Sound | | 33 USC 401 et seq.; 33 CFR 320–330 | Prohibits the unauthorized obstruction or alteration of any navigable water. Establishes requirements for structures or work in, above, or under navigable waters. | ARAR for remedial actions in Puget Sound. |

Table 3-3 Final RI/FS Work Plan Page 3 of 5

Table 3-3 – Potential ARARs, Action-SpecificBremerton Gas Works Superfund Site

| Remedial Activity | Act/Authority | Criteria/Issue | Citation | Brief Description | Applicability/Appropriateness |
|---|---|--------------------------------------|--|--|---|
| Dredging, Capping, and/or Discharge to Puget Sound (Continued) | Washington Hydraulics Code | Filling in Puget Sound | Chapter 75.20 and 77.55 RCW; Chapter 220-110 WAC | Establishes requirements for performing work that would use, divert, obstruct, or change the natural flow or bed of Puget Sound. | ARAR for shoreline excavation, dredging, and/or capping actions. Remedial actions must result in no net loss of aquatic habitat or function after sequential consideration of avoidance and mitigation. |
| | Federal Clean Air Act; Washington Clean Air Act; Puget Sound Air Clean Air Agency Regulations | Air Emission Discharges | 42 USC 7401 et seq.; Chapter 70.94 RCW; Chapter 173-400 WAC; PSCAA Regulation III | Regulates air emission discharges. | ARAR for remedial activities that generate fugitive dust or other air emissions, including treatment operations. |
| | Washington State Minimum Standards for Construction and Maintenance of Wells | Well Construction | Chapter 18.104 RCW; Chapter 173-160 WAC | Establishes minimum standards for the construction and decommissioning of all wells in the state of Washington. | ARAR for remedial activities that include installation and construction of monitoring wells or remediation wells used to inject any substance to remediate or control contamination. |
| Other Remedial Activities | Historic Preservation Act; Washington Historical Activities Act | Alteration of Historic Properties | 16 USC 470 et seq.; 36 CFR 800; Chapter 27 RCW | Requires the identification of historic properties potentially affected by remedial actions, and ways to avoid, minimize, or mitigate such effects. Historic property is any district, site, building, structure, or object included in or eligible for the National Register of Historic Places, including artifacts, records, and material remains related to such a property. | ARAR if historic properties are affected by remedial activities. No historic properties have been identified at the Site to date but could potentially be identified during remedial design. |
| | Archeological and Historic Preservation Act Alteration of Historic and Archaeological Properties | | 16 USC 469a-1 | Provides for the preservation of historical and archeological data that may be irreparably lost as a result of a federally approved project and mandates only preservation of the data. | ARAR if historical and archeological resources may be irreparably lost by implementation of remedial activities. |

Table 3-3 – Potential ARARs, Action-Specific

Bremerton Gas Works Superfund Site

Bremerton, Washington

| Remedial Activity | Act/Authority | Criteria/Issue | Citation | Brief Description | Applicability/Appropriateness |
|---|--|----------------------------------|--------------------------------|---|---|
| Other Remedial Activities (Continued) | Native American Graves Protection and Reparation Act | Alteration of American Graves | 25 USC 3001–3013; 43 CFR 10 | Requires federal agencies and museums that have possession of or control over Native American cultural items (including human remains, associated and unassociated funerary items, sacred objects, and objects of cultural patrimony) to compile an inventory of such items. Prescribes when such federal agencies and museums must return Native American cultural items. "Museums" are defined as any institution or state or local government agency that receives federal funds and has possession of, or control over, Native American cultural items. | ARAR if Native American cultural items are present in an excavation or dredging area. |

Notes:

ARAR = applicable or relevant and appropriate requirements

BMC = Bremerton Municipal Code

DNAPL = dense non-aqueous phase liquid

EPA = U.S. Environmental Protection Agency

FWHCA = Fish and Wildlife Habitat Conservation Area

MGP = manufactured gas plant

PSCCA = Puget Sound Clean Air Agency

RCW = Revised Code of Washington

SMP = Shoreline Master Program

TBC = to be considered

USC = United States Code

WAC = Washington Administrative Code

| | | EPA Ecological Soil Screening Levels - Birds | EPA Ecological Soil Screening Levels - Invertebrates | EPA Ecological Soil Screening Levels - Mammals | EPA Ecological Soil Screening Levels - Plants | EPA Regional Screening Levels (RSLs) - Residential Soil | EPA Regional Screening Levels (RSLs) - Industrial Soil | Laboratory MRL | 1 | s used for Data reening |
|--|------------|--|---|--|---|--|---|-------------------|-----------------------------|-------------------------------|
| Analyte | CAS Number | EPA, 2010 | EPA, 2010 | EPA, 2010 | EPA, 2010 | EPA, 2016 | EPA, 2016 | ARI, 2015 | Surface Soil (0-10 feet) | Subsurface Soil (>10 feet) |
| Conventionals (mg/kg) | | | | | | | | | | |
| Cyanide, WAD | 57-12-5 | | × | | | 2.3 | 15 | 0.05 | 2.3 | 2.3 |
| Cyanide, total | 57-12-5 | | | | | 2.3 | 15 | 0.05 | 2.3 | 2.3 |
| Metals (mg/kg) | | | | | | | | | | |
| Aluminum | 7429-90-5 | | France - | 122 | | 7,700 | 110,000 | | 7,700 | 7,700 |
| Antimony | 7440-36-0 | | 78 | 0.27 | | 3.1 | 47 | 0.2 | 0.27 | 3.1 |
| Arsenic | 7440-38-2 | 43 | | 46 | 18 | 0.68 | 3 | 0.5 | 0.68 | 0.68 |
| Barium | 7440-39-3 | | 330 | 2,000 | 1==1 | 1,500 | 22,000 | | 330 | 1,500 |
| Beryllium | 7440-41-7 | | 40 | 21 | | 16 | 230 | 0.2 | 16 | 16 |
| Cadmium | 7440-43-9 | 0.77 | 140 | 0.36 | 32 | 7.1 | 98 | 0.1 | 0.36 | 7.1 |
| Chromium | 7440-47-3 | 26 | (Appeller | 34 | | - | | 0.5 | 26 | == |
| Chromium III | 16065-83-1 | 26 | 1 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 | 34 | r an a | 12,000 | 180,000 | | 26 | 12,000 |
| Chromium VI | 18540-29-9 | - | - 10-10-10 | 130 | (==) | 0.3 | 6.3 | 0.4 | 0.3 | 0.3 |
| Cobalt | 7440-48-4 | 120 | - | 230 | 13 | 2.3 | 35 | 0.2 | 2.3 | 2.3 |
| Copper | 7440-50-8 | 28 | 80 | 49 | 70 | 310 | 4700 | 0.5 | 28 | 310 |
| Iron | 7439-89-6 | | 1 | | | 5,500 | 82,000 | | 5,500 | 5,500 |
| Lead | 7439-92-1 | 11 | 1,700 | 56 | 120 | 400 | 800 | 0.1 | 11 | 400 |
| Manganese | 7439-96-5 | 4,300 | 450 | 4,000 | 220 | 180 | 2,600 | 0.5 | 180 | 180 |
| Mercury | 7439-97-6 | | :: | | | 1.1 | 4.6 | 0.025 | 1.1 | 1.1 |
| Nickel | 7440-02-0 | 210 | 280 | 130 | 38 | 150 | 2,200 | 0.5 | 38 | 150 |
| Selenium | 7782-49-2 | 1.2 | 4.1 | 0.63 | 0.52 | 39 | 580 | 0.5 | 0.52 | 39 |
| Silver | 7440-22-4 | 4.2 | | 14 | 560 | 39 | 580 | 0.2 | 4.2 | 39 |
| Thallium | 7440-28-0 | | - | - | 186 | 0.078 | 1.2 | 0.2 | 0.078 | 0.078 |
| Vanadium | 7440-62-2 | 7.8 | 1 | 280 | - | 39 | 580 | | 7.8 | 39 |
| Zinc | 7440-66-6 | 46 | 120 | 79 | 160 | 2,300 | 35,000 | 4 | 46 | 2,300 |
| Metals, Organic (mg/kg) | | | | | | | | | | |
| Tributyltin | 688-73-3 | 122 | , lett | - | - | 2.3 | 35 | | 2.3 | 2.3 |
| Volatile Organic Compounds (VOCs) (mg/kg) | | | | | | *** | | | | |
| 1,1,1,2-Tetrachloroethane | 630-20-6 | | a== | : | | 2 | 8.8 | 0.001 | 2 | 2 |
| 1,1,1-Trichloroethane | 71-55-6 | | | 1.55 | | 810 | 3,600 | 0.001 | 810 | 810 |
| 1,1,2,2-Tetrachloroethane | 79-34-5 | 1 | CHIE | HH | (##) | 0.6 | 2.7 | 0.001 | 0.6 | 0.6 |
| 1,1,2-Trichloroethane | 79-00-5 | - | Total Control | | HH | 0.15 | 0.63 | 0.001 | 0.15 | 0.15 |
| 1,1,2-Trichlorotrifluoroethane (Freon 113) | 76-13-1 | | | 188 | (HH) | 4,000 | 17,000 | 0.002 | 4,000 | 4,000 |
| 1,1-Dichloroethane | 75-34-3 | | France - | 100 | | 3.6 | 16 | 0.001 | 3.6 | 3.6 |
| 1,1-Dichloroethene | 75-35-4 | | II | | | 23 | 100 | 0.001 | 23 | 23 |
| 1,2,3-Trichlorobenzene | 87-61-6 | | | | | 6.3 | 93 | 0.005 | 6.3 | 6.3 |
| 1,2,3-Trichloropropane | 96-18-4 | | | | | 0.0051 | 0.11 | 0.002 | 0.0051 | 0.0051 |
| 1,2,4-Trimethylbenzene | 95-63-6 | | : | | | 5.8 | 24 | 0.001 | 5.8 | 5.8 |
| 1,2-Dibromo-3-chloropropane | 96-12-8 | | ii | 1 | | 0.0053 | 0.064 | 0.005 | 0.0053 | 0.0053 |

| | | EPA Ecological Soil Screening Levels - Birds | EPA Ecological Soil Screening Levels - Invertebrates | EPA Ecological Soil Screening Levels - Mammals | EPA Ecological Soil Screening Levels - Plants | EPA Regional Screening Levels (RSLs) - Residential Soil | EPA Regional Screening Levels (RSLs) - Industrial Soil | Laboratory MRL | 1 | s used for Data reening |
|--|------------|--|---|--|---|--|---|-------------------|-----------------------------|-------------------------------|
| Analyte | CAS Number | EPA, 2010 | EPA, 2010 | EPA, 2010 | EPA, 2010 | EPA, 2016 | EPA, 2016 | ARI, 2015 | Surface Soil (0-10 feet) | Subsurface Soil (>10 feet) |
| Volatile Organic Compounds (VOCs) (mg/kg) (continued | | | | | | | | | | |
| 1,2-Dichloroethane | 107-06-2 | V | 100 | 100 | | 0.46 | 2 | 0.001 | 0.46 | 0.46 |
| 1,2-Dichloroethene, cis- | 156-59-2 | | - | N== | | 16 | 230 | 0.001 | 16 | 16 |
| 1,2-Dichloroethene, trans- | 156-60-5 | | | | | 160 | 2300 | 0.001 | 160 | 160 |
| 1,2-Dichloropropane | 78-87-5 | | : (| | : | 1 | 4.4 | 0.001 | 1 | 1 |
| 1,3,5-Trimethylbenzene (Mesitylene) | 108-67-8 | | - | | | 78 | 1,200 | 0.001 | 78 | 78 |
| 1,3-Dichloropropane | 142-28-9 | ·— | 2 | : | | 160 | 2,300 | 0.001 | 160 | 160 |
| 1,3-Dichloropropene, cis- | 10061-01-5 | | : | | | | | 0.001 | | |
| 1,3-Dichloropropene, trans- | 10061-02-6 | (Control | THE | EE | | N-19-40 | H | 0.001 | | |
| 1,4-Dichloro-2-butene, trans- | 110-57-6 | (200 | | 122 | (20) | 0.0074 | 0.032 | 0.005 | 0.0074 | 0.0074 |
| 2-Butanone (MEK) | 78-93-3 | | () () () () () () () () () () | | (200 1) | 2,700 | 19,000 | 0.005 | 2,700 | 2,700 |
| 2-Chlorotoluene | 95-49-8 | 19 <u>11111</u> | | | | 160 | 2,300 | | 160 | 160 |
| 2-Hexanone (Methyl butyl ketone) | 591-78-6 | | rame. | | | 20 | 130 | 0.005 | 20 | 20 |
| 4-Chlorotoluene | 106-43-4 | 7 | | | | 160 | 2,300 | 0.001 | 160 | 160 |
| 4-Isopropyltoluene (4-Cymene) | 99-87-6 | | | | | | | 0.001 | | |
| Acetone | 67-64-1 | | :1 | · | : | 6,100 | 67,000 | 0.005 | 6,100 | 6,100 |
| Acrolein | 107-02-8 | · | : 1,—— | :== | | 0.014 | 0.06 | 0.05 | 0.014 | 0.014 |
| Acrylonitrile | 107-13-1 | Q | : 2 | :== | | 0.25 | 1.1 | 0.005 | 0.25 | 0.25 |
| Benzene | 71-43-2 | | | | (ECC) | 1.2 | 5.1 | 0.001 | 1.2 | 1.2 |
| Bromobenzene | 108-86-1 | | Service Service | iee. | 4- 0 | 29 | 180 | 0.001 | 29 | 29 |
| Bromochloromethane | 74-97-5 | - | | 188 | (He) | 15 | 63 | 0.001 | 15 | 15 |
| Bromodichloromethane | 75-27-4 | - | 122 | THE | (800) | 0.29 | 1.3 | 0.001 | 0.29 | 0.29 |
| Bromoform (Tribromomethane) | 75-25-2 | | 100 | | | 19 | 86 | 0.001 | 19 | 19 |
| Bromomethane (Methyl bromide) | 74-83-9 | | 1 | | | 0.68 | 3 | 0.001 | 0.68 | 0.68 |
| Carbon disulfide | 75-15-0 | | | | | 77 | 350 | 0.001 | 77 | 77 |
| Carbon tetrachloride (Tetrachloromethane) | 56-23-5 | | : | | | 0.65 | 2.9 | 0.001 | 0.65 | 0.65 |
| Chlorobenzene | 108-90-7 | | :I== | :1== | - | 28 | 130 | 0.001 | 28 | 28 |
| Chloroethane | 75-00-3 | 0 | | :2== | | 1,400 | 5,700 | 0.001 | 1,400 | 1,400 |
| Chloroform | 67-66-3 | - | : december 1 | : 45500 | | 0.32 | 1.4 | 0.001 | 0.32 | 0.32 |
| Chloromethane | 74-87-3 | / | : | 1.000 | | 11 | 46 | 0.001 | 11 | 11 |
| Cyclohexane | 110-82-7 | - | I M | - | = | 650 | 2,700 | | 650 | 650 |
| Dibromochloromethane | 124-48-1 | (atr. | i a# | | | 8.3 | 39 | 0.001 | 8.3 | 8.3 |
| Dibromomethane | 74-95-3 | | | | | 2.4 | 9.9 | 0.001 | 2.4 | 2.4 |
| Dichlorodifluoromethane | 75-71-8 | | - | _ | | 8.7 | 37 | 0.001 | 8.7 | 8.7 |
| Dichloromethane (Methylene chloride) | 75-09-2 | | | | | 35 | 320 | 0.002 | 35 | 35 |
| Ethylbenzene | 100-41-4 | | _ | | | 5.8 | 25 | 0.001 | 5.8 | 5.8 |
| Ethylene dibromide (1,2-Dibromoethane) | 106-93-4 | | | | | 0.036 | 0.16 | 0.001 | 0.036 | 0.036 |
| Hexachlorobutadiene (Hexachloro-1,3-butadiene) | 87-68-3 | | | | | 1.2 | 5.3 | 0.005 | 1.2 | 1.2 |

| | | EPA Ecological Soil Screening Levels - Birds | EPA Ecological Soil Screening Levels - Invertebrates | EPA Ecological Soil Screening Levels - Mammals | EPA Ecological Soil Screening Levels - Plants | EPA Regional Screening Levels (RSLs) - Residential Soil | EPA Regional Screening Levels (RSLs) - Industrial Soil | Laboratory MRL | | s used for Data reening |
|---|------------|--|---|--|---|--|---|-------------------|-----------------------------|-------------------------------|
| Analyte | CAS Number | EPA, 2010 | EPA, 2010 | EPA, 2010 | EPA, 2010 | EPA, 2016 | EPA, 2016 | ARI, 2015 | Surface Soil (0-10 feet) | Subsurface Soil (>10 feet) |
| Volatile Organic Compounds (VOCs) (mg/kg) (continued |) | | | | | | | | | |
| Isopropylbenzene (Cumene) | 98-82-8 | | : new | : Carro | | 190 | 990 | 0.001 | 190 | 190 |
| Methyl acetate | 79-20-9 | | | | | 7,800 | 120,000 | | 7,800 | 7,800 |
| Methyl iodide (lodomethane) | 74-88-4 | - | | | | | | 0.001 | | |
| Methyl isobutyl ketone (4-Methyl-2-pentanone or (MIBK)) | 108-10-1 | | :1== | :1== | 11 | 3,300 | 14,000 | 0.005 | 3,300 | 3,300 |
| Methyl tert-butyl ether (MTBE) | 1634-04-4 | - | :1== | :1==. | (I mm) | 47 | 210 | 0.001 | 47 | 47 |
| n-Butylbenzene | 104-51-8 | | 12 | 12== | | 390 | 5,800 | 0.001 | 390 | 390 |
| n-Propylbenzene | 103-65-1 | y ana | 1.00-0-0-0 | 18-14-1 | | 380 | 2,400 | 0.001 | 380 | 380 |
| o-Xylene | 95-47-6 | _ | I | | - | 65 | 280 | 0.001 | 65 | 65 |
| sec-Butylbenzene | 135-98-8 | _ | I | | - | 780 | 12,000 | 0.001 | 780 | 780 |
| Styrene | 100-42-5 | - | 1-4 | | - | 600 | 3,500 | 0.001 | 600 | 600 |
| tert-Butylbenzene | 98-06-6 | | | | | 780 | 12,000 | 0.001 | 780 | 780 |
| Tetrachloroethene (PCE) | 127-18-4 | | | | | 8.1 | 39 | 0.001 | 8.1 | 8.1 |
| Toluene | 108-88-3 | | × | | | 490 | 4,700 | 0.001 | 490 | 490 |
| Total xylene (reported, not calculated) | 1330-20-7 | | | | | 58 | 250 | 0.002 | 58 | 58 |
| Trichloroethene (TCE) | 79-01-6 | - | 1 | 1 | | 0.41 | 1.9 | 0.001 | 0.41 | 0.41 |
| Trichlorofluoromethane (Fluorotrichloromethane) | 75-69-4 | · | :3 | :== | | 2,300 | 35,000 | 0.001 | 2,300 | 2,300 |
| Vinyl acetate | 108-05-4 | 0 | :== | :2 | | 91 | 380 | 0.005 | 91 | 91 |
| Vinyl chloride | 75-01-4 | | : | : | | 0.059 | 1.7 | 0.001 | 0.059 | 0.059 |
| Semivolatile Organic Componds (SVOCs) (mg/kg) | • | | | | • | | | | • | |
| 1,2,4,5-Tetrachlorobenzene | 95-94-3 | · | 0 | n== | | 2.3 | 35 | 0.067 | 2.3 | 2.3 |
| 1,2,4-Trichlorobenzene | 120-82-1 | · | n | n | | 5.8 | 26 | 0.067 | 5.8 | 5.8 |
| 1,2-Dichlorobenzene | 95-50-1 | | - | | | 180 | 930 | 0.067 | 180 | 180 |
| 1,3-Dichlorobenzene | 541-73-1 | _ | i | | - | | - | 0.067 | | |
| 1,4-Dichlorobenzene | 106-46-7 | - | [| 1 | - | 2.6 | 11 | 0.067 | 2.6 | 2.6 |
| 1,4-Dioxane | 123-91-1 | _ | | | - | 5.3 | 24 | 0.067 | 5.3 | 5.3 |
| 2,2'-Oxybis (1-chloropropane) | 108-60-1 | 1-m- | :: | :: - | | 310 | 4,700 | 0.067 | 310 | 310 |
| 2,3,4,6-Tetrachlorophenol | 58-90-2 | | - | - | - | 190 | 2,500 | 0.067 | 190 | 190 |
| 2,4,5-Trichlorophenol | 95-95-4 | | | | | 630 | 8,200 | 0.33 | 630 | 630 |
| 2,4,6-Trichlorophenol | 88-06-2 | | - | | | 6.3 | 82 | 0.33 | 6.3 | 6.3 |
| 2,4-Dichlorophenol | 120-83-2 | | :1== | : | () | 19 | 250 | 0.33 | 19 | 19 |
| 2,4-Dimethylphenol | 105-67-9 | | - | | : > | 130 | 1600 | 0.067 | 130 | 130 |
| 2,4-Dinitrophenol | 51-28-5 | | 1.55 | - | | 13 | 160 | 0.67 | 13 | 13 |
| 2,4-Dinitrotoluene | 121-14-2 | Y ONE | 10000 | | , mak | 1.7 | 7.4 | 0.33 | 1.7 | 1.7 |
| 2,6-Dinitrotoluene | 606-20-2 | _ | (((((((((((((((((((| (1000) (1000) | (UNI) | 0.36 | 1.5 | 0.33 | 0.36 | 0.36 |
| 2-Chloronaphthalene | 91-58-7 | (| | == | l a- ti | 480 | 6,000 | 0.067 | 480 | 480 |

| , | | EPA Ecological Soil Screening Levels - Birds | EPA Ecological Soil Screening Levels - Invertebrates | EPA Ecological Soil Screening Levels - Mammals | EPA Ecological Soil Screening Levels - Plants | EPA Regional Screening Levels (RSLs) - Residential Soil | EPA Regional Screening Levels (RSLs) - Industrial Soil | Laboratory MRL | | s used for Data reening |
|---|------------|--|---|--|--|--|---|-------------------|-----------------------------|-------------------------------|
| Analyte | CAS Number | EPA, 2010 | EPA, 2010 | EPA, 2010 | EPA, 2010 | EPA, 2016 | EPA, 2016 | ARI, 2015 | Surface Soil (0-10 feet) | Subsurface Soil (>10 feet) |
| Semivolatile Organic Componds (SVOCs) (mg/kg) (contin | | | | | | | | | _ | |
| 2-Chlorophenol | 95-57-8 | | 1900 | | | 39 | 580 | 0.067 | 39 | 39 |
| 2-Methylphenol (o-Cresol) | 95-48-7 | | · | | | 320 | 4,100 | 0.067 | 320 | 320 |
| 2-Nitroaniline | 88-74-4 | | | | | 63 | 800 | 0.33 | 63 | 63 |
| 2-Nitrophenol | 88-75-5 | | 1 | | | - | | 0.067 | | == |
| 3,3'-Dichlorobenzidine | 91-94-1 | | | | | 1.2 | 5.1 | 0.33 | 1.2 | 1.2 |
| 3-Methylphenol & 4-Methylphenol (m&p-Cresol) | 1319-77-3 | | × | | | 630 | 8200 | | 630 | 630 |
| B-Methylphenol (m-Cresol) | 108-39-4 | | | | | 320 | 4100 | | 320 | 320 |
| 3-Nitroaniline | 99-09-2 | Control | | THE STATE OF THE S | THE STATE OF THE S | | 144 | 0.33 | | |
| 4-Bromophenyl-phenyl ether | 101-55-3 | ASSES. | 1 | 18-30- | 1 311 2 | | 100 | 0.067 | | |
| 1-Chloro-3-methylphenol | 59-50-7 | (ledin) | 188 | (EE | .55 | 630 | 8,200 | 0.33 | 630 | 630 |
| 4-Chloroaniline | 106-47-8 | - | - | | | 2.7 | 11 | 0.33 | 2.7 | 2.7 |
| 4-Methylphenol (p-Cresol) | 106-44-5 | | × | | | 630 | 8200 | 0.067 | 630 | 630 |
| 1-Nitroaniline | 100-01-6 | | - | | | 25 | 110 | 0.33 | 25 | 25 |
| 1-Nitrophenol | 100-02-7 | | - | | | | | 0.33 | | |
| Aniline | 62-53-3 | | 1 | | | 44 | 400 | 0.067 | 44 | 44 |
| Benzidine | 92-87-5 | | :== | : | | 0.00053 | 0.01 | | 0.00053 | 0.00053 |
| Benzoic acid | 65-85-0 | | : 2. | : | | 25,000 | 330,000 | 0.67 | 25,000 | 25,000 |
| Benzyl alcohol | 100-51-6 | | | · · | | 630 | 8,200 | 0.33 | 630 | 630 |
| Biphenyl (1,1'-Biphenyl) | 92-52-4 | | 10-12-20 10-12-20 | 1 (min 22 m) | - | 4.7 | 20 | 0.005 | 4.7 | 4.7 |
| pis(2-Chloroethoxy)methane | 111-91-1 | CORECTOR CONTRACTOR CO | I E # | T age | | 19 | 250 | 0.067 | 19 | 19 |
| pis(2-Chloroethyl)ether | 111-44-4 | (uzu | i ax | | - | 0.23 | 1 | 0.067 | 0.23 | 0.23 |
| pis(2-Ethylhexyl)phthalate | 117-81-7 | | (Care) | | | 39 | 160 | 0.067 | 39 | 39 |
| Butylbenzyl phthalate | 85-68-7 | | r un | | | 290 | 1,200 | 0.067 | 290 | 290 |
| Dibenzofuran | 132-64-9 | | | | | 7.3 | 100 | 0.067 | 7.3 | 7.3 |
| Diethyl phthalate | 84-66-2 | | | | | 5,100 | 66,000 | 0.067 | 5,100 | 5,100 |
| Dimethyl phthalate | 131-11-3 | | | | | | | 0.067 | | |
| Di-n-butyl phthalate | 84-74-2 | | : | : | | 630 | 8,200 | 0.067 | 630 | 630 |
| Dinitro-o-cresol (4,6-Dinitro-2-methylphenol) | 534-52-1 | , | | | | 0.51 | 6.6 | 0.67 | 0.51 | 0.51 |
| Di-n-octyl phthalate | 117-84-0 | <u></u> | | | | 63 | 820 | 0.067 | 63 | 63 |
| Hexachlorobenzene | 118-74-1 | <u>-</u> | | | | 0.21 | 0.96 | 0.067 | 0.21 | 0.21 |
| Hexachlorocyclopentadiene | 77-47-4 | - | 120/320 | | - | 0.18 | 0.75 | 0.33 | 0.18 | 0.18 |
| Hexachloroethane | 67-72-1 | | | 100 | ### | 1.8 | 8 | 0.067 | 1.8 | 1.8 |
| sophorone | 78-59-1 | NAME: | | | | 570 | 2,400 | 0.067 | 570 | 570 |
| Vitrobenzene | 98-95-3 | | \ <u>-</u> | | | 5.1 | 22 | 0.067 | 5.1 | 5.1 |
| n-Nitrosodimethylamine | 62-75-9 | | | | | 0.002 | 0.034 | 0.007 | 0.002 | |
| n-Nitrosodi-n-propylamine | 621-64-7 | | | | | 0.002 | 0.33 | 0.067 | 0.002 | 0.002 0.078 |

Bremerton, Washington

| | | EPA Ecological Soil Screening Levels - Birds | EPA Ecological Soil Screening Levels - Invertebrates | EPA Ecological Soil Screening Levels - Mammals | EPA Ecological Soil Screening Levels - Plants | | EPA Regional Screening Levels (RSLs) - Industrial Soil | Laboratory MRL | | s used for Data eening | | |
|---|---------------|--|---|--|---|---------------------|---|-------------------|-----------------------------|-------------------------------|--|--|
| Analyte | CAS Number | EPA, 2010 | EPA, 2010 | EPA, 2010 | EPA, 2010 | EPA, 2016 | EPA, 2016 | ARI, 2015 | Surface Soil (0-10 feet) | Subsurface Soil (>10 feet) | | |
| Semivolatile Organic Componds (SVOCs) (mg/kg) (continued) | | | | | | | | | | | | |
| n-Nitrosodiphenylamine | 86-30-6 | - | - | | | 110 | 470 | 0.067 | 110 | 110 | | |
| Pentachlorophenol | 87-86-5 | 2.1 | 31 | 2.8 | 5 | 1 | 4 | 0.33 | 1 | 1 | | |
| Phenol | 108-95-2 | | - | - | - | 1900 | 25000 | 0.067 | 1900 | 1900 | | |
| Polycyclic Aromatic Hydrocarbons (PAHs) (mg/kg) | | | | | | | | | | | | |
| 1-Methylnaphthalene | 90-12-0 | | | | | 18 | 73 | 0.005 | 18 | 18 | | |
| 2-Methylnaphthalene | 91-57-6 | | « | | | 24 | 300 | 0.005 | 24 | 24 | | |
| Acenaphthene | 83-32-9 | | - | | | 360 | 4,500 | 0.005 | 360 | 360 | | |
| Acenaphthylene | 208-96-8 | | | : | | _ | | 0.005 | | | | |
| Anthracene | 120-12-7 | | | · | | 1,800 | 23,000 | 0.005 | 1,800 | 1,800 | | |
| Benzo(a)anthracene | 56-55-3 | - | \ | | , pos t | 0.16 | 2.9 | 0.005 | 0.16 | 0.16 | | |
| Benzo(a)pyrene | 50-32-8 | , | < | ~ | | 0.016 | 0.29 | 0.005 | 0.016 | 0.016 | | |
| Benzo(b)fluoranthene | 205-99-2 | | - | :== | *** | 0.16 | 2.9 | 0.005 | 0.16 | 0.16 | | |
| Benzo(g,h,i)perylene | 191-24-2 | | | 1 | - | | | 0.005 | | | | |
| Benzo(k)fluoranthene | 207-08-9 | V ote | 1 =1 | S | | 1.6 | 29 | 0.005 | 1.6 | 1.6 | | |
| Chrysene | 218-01-9 | | | | | 16 | 290 | 0.005 | 16 | 16 | | |
| Dibenzo(a,h)anthracene | 53-70-3 | | \ | · | | 0.016 | 0.29 | 0.005 | 0.016 | 0.016 | | |
| Fluoranthene | 206-44-0 | | × | · | | 240 | 3,000 | 0.005 | 240 | 240 | | |
| Fluorene | 86-73-7 | | : | | | 240 | 3,000 | 0.005 | 240 | 240 | | |
| Indeno(1,2,3-c,d)pyrene | 193-39-5 | - | | | | 0.16 | 2.9 | 0.005 | 0.16 | 0.16 | | |
| Naphthalene | 91-20-3 | | - N ewson | · | (200) | 3.8 | 17 | 0.005 | 3.8 | 3.8 | | |
| Phenanthrene | 85-01-8 | | N ame | · | . | N atio k | | 0.005 | | | | |
| Pyrene | 129-00-0 | v o | | | A.A. | 180 | 2,300 | 0.005 | 180 | 180 | | |
| Total HPAH | : 2-1-2 | (*** | 18 | 1.1 | | 122 | - | | 1.1 | | | |
| Total LPAH | 1000 | · m | 29 | 100 | 10 | V ala ti | | | 29 | | | |
| Total PAH | · | ale | 1 | 5 1.0 | | U nio | | | | | | |

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Bremerton, Washington

| | | EPA Ecological Soil Screening Levels - Birds | EPA Ecological Soil Screening Levels - Invertebrates | EPA Ecological Soil Screening Levels - Mammals | EPA Ecological Soil Screening Levels - Plants | EPA Regional Screening Levels (RSLs) - Residential Soil | EPA Regional Screening Levels (RSLs) - Industrial Soil | Laboratory MRL | 1 | s used for Data reening |
|--|------------|--|---|--|---|--|---|-------------------|-----------------------------|-------------------------------|
| Analyte | CAS Number | EPA, 2010 | EPA, 2010 | EPA, 2010 | EPA, 2010 | EPA, 2016 | EPA, 2016 | ARI, 2015 | Surface Soil (0-10 feet) | Subsurface Soil (>10 feet) |
| Organochlorine Pesticides (mg/kg) | | | | | | | | | • | |
| Hexachlorocyclohexane, alpha-BHC | 319-84-6 | | | | | 0.086 | 0.36 | 0.0017 | 0.086 | 0.086 |
| Hexachlorocyclohexane, beta-BHC | 319-85-7 | | | | | 0.3 | 1.3 | 0.0017 | 0.3 | 0.3 |
| Hexachlorocyclohexane, gamma-BHC (Lindane) | 58-89-9 | | | | | 0.57 | 2.5 | 0.0017 | 0.57 | 0.57 |
| Hexachlorocyclohexane, delta-BHC | 319-86-8 | | | | | | | 0.0017 | | |
| Heptachlor | 76-44-8 | | | | | 0.13 | 0.63 | 0.0017 | 0.13 | 0.13 |
| Aldrin | 309-00-2 | | - | | | 0.039 | 0.18 | 0.0017 | 0.039 | 0.039 |
| Heptachlor epoxide | 1024-57-3 | | | | | 0.07 | 0.33 | 0.0017 | 0.07 | 0.07 |
| Chlordane | 57-74-9 | | | | | | | 0.0017 | | |
| Chlordane (technical) | 12789-03-6 | | | | | 1.7 | 7.7 | 0.0017 | 1.7 | 1.7 |
| Endosulfan-alpha (I) | 959-98-8 | | | | | | | 0.0017 | | |
| 4,4'-DDE (p,p'-DDE) | 72-55-9 | | | | | 2 | 9.3 | 0.0033 | 2 | 2 |
| Dieldrin | 60-57-1 | 0.022 | | 0.0049 | | 0.034 | 0.14 | 0.0033 | 0.0049 | 0.034 |
| Endrin | 72-20-8 | | | | | 1.9 | 25 | 0.0033 | 1.9 | 1.9 |
| Endosulfan-beta (II) | 33213-65-9 | | | | | | | 0.0033 | | |
| 4,4'-DDD (p,p'-DDD) | 72-54-8 | | | | | 2.3 | 9.6 | 0.0033 | 2.3 | 2.3 |
| Endrin aldehyde | 7421-93-4 | | | | | | | 0.0033 | | |
| 4,4'-DDT (p,p'-DDT) | 50-29-3 | | | | | 1.9 | 8.5 | 0.0033 | 1.9 | 1.9 |
| Endosulfan sulfate | 1031-07-8 | | | | | | | 0.0033 | | |
| Methoxychlor | 72-43-5 | | | | | 32 | 410 | 0.0017 | 32 | 32 |
| Polychlorinated Biphenyls (PCBs) (mg/kg) | | • | | • | | | | | • | |
| Aroclor 1016 | 12674-11-2 | | | | | 0.41 | 5.1 | 0.33 | 0.41 | 0.41 |
| Aroclor 1221 | 11104-28-2 | | | | | 0.2 | 0.83 | 0.33 | 0.2 | 0.2 |
| Aroclor 1232 | 11141-16-5 | | | | | 0.17 | 0.72 | 0.33 | 0.17 | 0.17 |
| Aroclor 1242 | 53469-21-9 | | | | | 0.23 | 0.95 | 0.33 | 0.23 | 0.23 |
| Aroclor 1248 | 12672-29-6 | | | | | 0.23 | 0.95 | 0.33 | 0.23 | 0.23 |
| Aroclor 1254 | 11097-69-1 | | | | | 0.12 | 0.97 | 0.33 | 0.12 | 0.12 |
| Aroclor 1260 | 11096-82-5 | | | | | 0.24 | 0.99 | 0.33 | 0.24 | 0.24 |
| Total PCB Aroclors | 1336-36-3 | | | | | 0.23 | 0.94 | 0.33 | 0.23 | 0.23 |

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Table 3-4 - Development of Initial PRGs for Soil

Bremerton Gas Works Superfund Site Bremerton, Washington

| | | EPA Ecological Soil Screening Levels - Birds | EPA Ecological Soil Screening Levels - Invertebrates | | EPA Ecological Soil Screening Levels - Plants | Levels (RSLs) - | (A) | Laboratory MRL | The state of the s | s used for Data reening |
|---------|------------|--|---|-----------|---|-----------------|-----------|-------------------|--|-------------------------------|
| Analyte | CAS Number | EPA, 2010 | EPA, 2010 | EPA, 2010 | EPA, 2010 | EPA, 2016 | EPA, 2016 | ARI, 2015 | Surface Soil (0-10 feet) | Subsurface Soil (>10 feet) |

Notes

Compounds frequently associated with MGP-operations.

- indicates not available

CAS = Chemical Abstract Services

EPA = U.S. Environmental Protection Agency

HPAH = high molecular weight PAH

LPAH = low molecular weight PAH

kg = kilogram

mg = miligram

MGP = manufactured gas plant

ng = nanogram

PRG = preliminary remediation goal

RCRA = Resource Conservation and Recovery Act

RSL = regional screening level

ug = microgram

WAD = Weak Acid Dissociable Cyanide

References:

EPA, 2010. Ecological Soil Screening Levels. Updated October 20, 2010. Cited: January 15, 2014. Available from: http://www.epa.gov/ecotox/ecossl/.

EPA, 2016. EPA Regional Screening Levels. May 2016. Available from: https://www.epa.gov/risk/regional-screening-levels-rsls-generic-tables-may-2016.

Table 3-5 - Development of Initial PRGs for Groundwater Bremerton Gas Works Superfund Site Bremerton, Washington

| Analista | CAS Number | EPA Regional Screening Levels (RSLs) - MCL EPA, 2016 | EPA Regional Screening Levels (RSLs) - Tapwater EPA, 2016 | Laboratory MRL ARI, 2015 | Initial PRGs used for Data |
|--|------------------------|---|--|-----------------------------|-------------------------------|
| Analyte Conventionals (mg/L) | CAS Number | EPA, 2016 | EPA, 2016 | ARI, 2015 | Screening |
| Cyanide, free | 57-12-5 | 0.2 | 0.15 | 0.00500 | 0.15 |
| Metals (ug/L) | 0. 120 | 0.000 | Special Petronic | | October Process |
| Antimony | 7440-36-0 | 6 | 0.78 | 0.2 | 0.78 |
| Arsenic | 7440-38-2 | 10 | 0.052 | 0.2 | 0.052 |
| Barium | 7440-39-3 | 2,000 | 380 | | 380 |
| Beryllium | 7440-41-7 | 4 | 2.5 | 0.2 | 2.5 |
| Cadmium | 7440-43-9 | 5 | 0.92 | 0.1 | 0.92 |
| Chromium | 7440-47-3 | 100 | 1 | 0.5 | 100 |
| Chromium III | 16065-83-1 | (m-12) | 2,200 | n/a | 2,200 |
| Chromium VI | 18540-29-9 | - | 0.035 | 0.01 | 0.035 |
| Cobalt | 7440-48-4 | _ | 0.6 | | 0.6 |
| Copper | 7440-50-8 | 1,300 | 80 | 0.5 | 80 |
| Lead | 7439-92-1 | 15 | 15 | 0.1 | 15 |
| Manganese | 7439-96-5 | - | 43 | 0.400 | 43 |
| Mercury | 7439-97-6 | 2 | 0.063 | 0.100 | 0.063 |
| Nickel Selenium | 7440-02-0 7782-49-2 | 50 | 39 10 | 0.5 0.5 | 39 10 |
| Selenium | 7/82-49-2 | 100000000 | 9.4 | 0.5 | 9.4 |
| Silver | 7440-22-4 | 2 | 9.4 0.02 | 0.2 | 9.4 0.02 |
| i nailium Vanadium | 7440-28-0 | | 8.6 | 0.∠ | 8.6 |
| Zinc | 7440-62-2 | | 6.0 | 4 | 600 |
| Metals, Olrganic (ug/L) | 7440-00-0 | | 000 | 7 | 000 |
| Tributyltin | 688-73-3 | <u> </u> | 0.37 | n/a | 0.37 |
| Volatile Organic Compounds (VOCs) (ug/L) | 000 70 0 | | 0.07 | Tira | 0.07 |
| 1,1,1,2-Tetrachloroethane | 630-20-6 | | 0.57 | 0.200 | 0.57 |
| 1,1,1-Trichloroethane | 71-55-6 | 200 | 800 | 0.200 | 200 |
| 1,1,2,2-Tetrachloroethane | 79-34-5 | - | 0.076 | 0.200 | 0.076 |
| 1,1,2-Trichloroethane | 79-00-5 | 5 | 0.041 | 0.200 | 0.041 |
| 1,1,2-Trichlorotrifluoroethane (Freon 113) | 76-13-1 | | 5,500 | 0.200 | 5,500 |
| 1,1-Dichloroethane | 75-34-3 | 10-10-0 0-10-0 | 2.8 | 0.200 | 2.8 |
| 1,1-Dichloroethene | 75-35-4 | 7 | 28 | 0.200 | 7 |
| 1,2,3-Trichlorobenzene | 87-61-6 | - | 0.7 | 0.500 | 0.7 |
| 1,2,3-Trichloropropane | 96-18-4 | | 0.00075 | 0.500 | 0.00075 |
| 1,2,4-Trimethylbenzene | 95-63-6 | | 1.5 | 0.200 | 1.5 |
| 1,2-Dibromo-3-chloropropane | 96-12-8 | 0.2 | 0.00033 | 0.500 | 0.00033 |
| 1,2-Dichloroethane | 107-06-2 | 5 | 0.17 | 0.200 | 0.17 |
| 1,2-Dichloroethene, cis- | 156-59-2 | 70 | 3.6 | 0.200 | 3.6 |
| 1,2-Dichloroethene, trans- | 156-60-5 | 100 | 36 | 0.200 | 36 |
| 1,2-Dichloropropane | 78-87-5 | 5 | 0.44 | 0.200 | 0.44 |
| 1,3,5-Trimethylbenzene (Mesitylene) | 108-67-8 | _ | 12 | 0.200 | 12 |
| 1,3-Dichloropropane | 142-28-9 | (| 37 | 0.200 | 37 |
| 1,3-Dichloropropene, cis- | 10061-01-5 | 1 -11 | 1== | 0.200 | |
| 1,3-Dichloropropene, trans- | 10061-02-6 | 1 | | 0.200 | |
| 1,4-Dichloro-2-butene, trans- | 110-57-6 | 7 | 0.0013 | 1.00 | 0.0013 |
| 2-Butanone (MEK) 2-Chlorotoluene | 78-93-3 95-49-8 | - | 560 24 | 5.00 | 560 24 |
| 2-Unioroloidene 2-Hexanone (Methyl butyl ketone) | 591-78-6 | 1 | 3.8 | 0.200 5.00 | 3.8 |
| 4-Chlorotoluene | 106-43-4 | | 25 | 0.200 | 25 |
| 4-Isopropyltoluene (4-Cymene) | 99-87-6 | - | 23 | 0.200 | 23 |
| Acetone | 67-64-1 | | 1,400 | 5.00 | 1,400 |
| Acrolein | 107-02-8 | | 0.0042 | 5.00 | 0.0042 |
| Acrylonitrile | 107-02-8 | | 0.052 | 1.00 | 0.052 |
| Benzene | 71-43-2 | 5 | 0.46 | 0.200 | 0.46 |
| Bromobenzene | 108-86-1 | _ | 6.2 | 0.200 | 6.2 |
| Bromochloromethane | 74-97-5 | | 8.3 | 0.200 | 8.3 |
| Bromodichloromethane | 75-27-4 | 80 | 0.13 | 0.200 | 0.13 |
| Bromoform (Tribromomethane) | 75-25-2 | 80 | 3.3 | 0.200 | 3.3 |
| Bromomethane (Methyl bromide) | 74-83-9 | | 0.75 | 1.00 | 0.75 |
| Carbon disulfide | 75-15-0 | - | 81 | 0.200 | 81 |
| Carbon tetrachloride (Tetrachloromethane) | 56-23-5 | 5 | 0.46 | 0.200 | 0.46 |
| Chlorobenzene | 108-90-7 | 100 | 7.8 | 0.200 | 7.8 |
| Chloroethane | 75-00-3 | 100 | 2,100 | 0.200 | 2,100 |
| Chloroform | 67-66-3 | 80 | 0.22 | 0.200 | 0.22 |
| Chloromethane | 74-87-3 | | 19 | 0.500 | 19 |
| | | + | | | 1 |
| Cyclohexane | 110-82-7 | | 1,300 | n/a | 1,300 |

Table 3-5 - Development of Initial PRGs for Groundwater Bremerton Gas Works Superfund Site Bremerton, Washington

| Analista | CAS Number | EPA Regional Screening Levels (RSLs) - MCL EPA, 2016 | EPA Regional Screening Levels (RSLs) - Tapwater EPA, 2016 | Laboratory MRL ARI, 2015 | Initial PRGs used for Data |
|---|------------|---|--|-----------------------------|-------------------------------|
| Analyte | CAS Number | LFA, 2010 | LFA, 2010 | AIN, 2015 | Screening |
| Volatile Organic Compounds (VOCs) (ug/L) (contin Dibromomethane | 74-95-3 | | 0.83 | 0.200 | 0.83 |
| Dichlorodifluoromethane | 75-71-8 | | 20 | 0.200 | 20 |
| Dichloromethane (Methylene chloride) | 75-09-2 | 5 | 11 | 1.00 | 5 |
| Ethylbenzene | 100-41-4 | 700 | 1.5 | 0.200 | 1.5 |
| Ethylene dibromide (1,2-Dibromoethane) | 106-93-4 | 0.05 | 0.0075 | 0.200 | 0.0075 |
| Hexachlorobutadiene (Hexachloro-1,3-butadiene) | 87-68-3 | (co | 0.14 | 0.500 | 0.14 |
| Isopropylbenzene (Cumene) | 98-82-8 | 14-15-0 (4-15-1) | 45 | 0.200 | 45 |
| Methyl acetate | 79-20-9 | | 2000 | n/a | 2000 |
| Methyl iodide (lodomethane) | 74-88-4 | | | 1.00 | ())1 |
| Methyl isobutyl ketone (4-Methyl-2-pentanone or (MIBK)) | 108-10-1 | | 630 | 5.00 | 630 |
| Methyl tert-butyl ether (MTBE) | 1634-04-4 | | 14 | 0.500 | 14 |
| n-Butylbenzene | 104-51-8 | (coo | 100 | 0.200 | 100 |
| n-Propylbenzene | 103-65-1 | (8-100) (8-100) | 66 | 0.200 | 66 |
| o-Xylene | 95-47-6 | (200) | 19 | 0.200 | 19 |
| sec-Butylbenzene | 135-98-8 | · | 200 | 0.200 | 200 |
| Styrene | 100-42-5 | 100 | 120 | 0.200 | 100 |
| tert-Butylbenzene | 98-06-6 | _ | 69 | 0.200 | 69 |
| Tetrachloroethene (PCE) | 127-18-4 | 5 | 4.1 | 0.200 | 4.1 |
| Toluene | 108-88-3 | 1000 | 110 | 0.200 | 110 |
| Total xylene (reported, not calculated) | 1330-20-7 | 10000 | 19 | n/a | 19 |
| Total Xylene | | | | n/a | |
| Trichloroethene (TCE) | 79-01-6 | 5 | 0.28 | 0.200 | 0.28 |
| Trichlorofluoromethane (Fluorotrichloromethane) | 75-69-4 | · | 520 | 0.200 | 520 |
| Vinyl acetate | 108-05-4 | - | 41 | 0.200 | 41 |
| Vinyl chloride | 75-01-4 | 2 | 0.019 | 0.200 | 0.019 |
| Semivolatile Organic Compounds (SVOCs) (ug/L) 1,2,4,5-Tetrachlorobenzene | 95-94-3 | | 0.17 | n/a | 0.17 |
| 1,2,4-Trichlorobenzene | 120-82-1 | 70 | 0.17 | 0.254 | 0.17 |
| 1,2-Dichlorobenzene | 95-50-1 | 600 | 30 | 0.250 | 30 |
| 1,3-Dichlorobenzene | 541-73-1 | | | 0.266 | |
| 1,4-Dichlorobenzene | 106-46-7 | 75 | 0.48 | 0267 | 0.48 |
| 1,4-Dioxane | 123-91-1 | | 0.46 | 0.4 | 0.46 |
| 2,2'-Oxybis (1-chloropropane) | 108-60-1 | | 71 | 0.241 | 71 |
| 2,3,4,6-Tetrachlorophenol | 58-90-2 | \ | 24 | 0.244 | 24 |
| 2,4,5-Trichlorophenol | 95-95-4 | | 120 | 1.10 | 120 |
| 2,4,6-Trichlorophenol | 88-06-2 | - | 1.2 | 1.04 | 1.2 |
| 2,4-Dichlorophenol | 120-83-2 | · | 4.6 | 1.11 | 4.6 |
| 2,4-Dimethylphenol | 105-67-9 | - | 36 | 1.12 | 36 |
| 2,4-Dinitrophenol | 51-28-5 | - | 3.9 | 3.35 | 3.9 |
| 2,4-Dinitrotoluene | 121-14-2 | | 0.24 | 1.12 | 0.24 |
| 2,6-Dinitrotoluene | 606-20-2 | | 0.049 | 1.14 | 0.049 |
| 2-Chloronaphthalene | 91-58-7 | (| 75 | 0.248 | 75 |
| 2-Chlorophenol | 95-57-8 | 10 - 17 - 17 - 17 - 17 - 17 - 17 - 17 - | 9.1 | 0.220 | 9.1 |
| 2-Methylphenol (o-Cresol) | 95-48-7 | · · · · · · | 93 | 0.211 | 93 |
| 2-Nitroaniline | 88-74-4 | | 19 | 1.46 | 19 |
| 2-Nitrophenol | 88-75-5 | (| 1 | 0.263 | |
| 3,3'-Dichlorobenzidine | 91-94-1 | = | 0.13 | 1.77 | 0.13 |
| 3-Methylphenol & 4-Methylphenol (m&p-Cresol) | 1319-77-3 | · | 150 | n/a | 150 |
| 3-Methylphenol (m-Cresol) | 108-39-4 | _ | 93 | n/a | 93 |
| 3-Nitroaniline | 99-09-2 | _ | | 1.53 | |
| 4-Bromophenyl-phenyl ether | 101-55-3 | | | 0.238 | |
| 4-Chloro-3-methylphenol | 59-50-7 | | 140 | 1.12 | 140 |
| 4-Chloroaniline | 106-47-8 | - | 0.37 | 1.73 | 0.37 |
| 4-Methylphenol (p-Cresol) | 106-44-5 | | 190 | 0.468 | 190 |
| 4-Nitroaniline | 100-01-6 | - | 3.8 | 2.02 | 3.8 |
| 4-Nitrophenol | 100-02-7 | - | | 1.75 | |
| Acetophenone | 98-86-2 | · | 190 | n/a | 190 |
| Aniline | 62-53-3 | - | 13 | 0.973 | 13 |
| Atrazine | 1912-24-9 | 3 | 0.3 | n/a | 0.3 |
| Benzaldehyde | 100-52-7 | - | 19 | n/a | 19 |
| Benzidine | 92-87-5 | | 0.00011 | n/a | 0.00011 |
| Benzoic acid | 65-85-0 | - | 7500 | 3.92 | 7500 |
| Benzyl alcohol | 100-51-6 | 100 | 200 | 0.552 | 200 |
| Biphenyl (1,1'-Biphenyl) | 92-52-4 | - | 0.083 | n/a | 0.083 |
| bis(2-Chloroethoxy)methane | 111-91-1 | · | 5.9 | 0.237 | 5.9 |

Table 3-5 - Development of Initial PRGs for Groundwater

Bremerton Gas Works Superfund Site

Bremerton, Washington

| Semivolatile Organic Compounds (SVOCs) (ug/L) (continued) bis(2-Chloroethyl)ether 111-44 bis(2-Ethylhexyl)phthalate 117-81 Butylbenzyl phthalate 85-68 Caprolactam 105-66 Caprolactam 132-64 Dibenzofuran 132-64 Diethyl phthalate 84-66 Dimethyl phthalate 131-11 Di-n-butyl phthalate 131-11 Di-n-butyl phthalate 84-74 Dinitro-o-cresol (4,6-Dinitro-2-methylphenol) 534-52 Di-n-octyl phthalate 117-84 Hexachlorobenzene 118-74 Hexachlorocyclopentadiene 77-47 Hexachlorocyclopentadiene 67-72 Isophorone 78-59 Nitrobenzene 98-95 n-Nitrosodimethylamine 62-75 n-Nitrosodimethylamine 62-75 n-Nitrosodimethylamine 621-64 n-Nitrosodiphenylamine 86-30 Phenol 108-95 Polycyclic Aromatic Hydrocarbons (PAHs)(ug/L) 1-Methylnaphthalene 90-12 2-Methylnaphthalene 90-12 2-Methylnaphthalene 120-12 Benzo(a)anthracene 120-12 Benzo(a)anthracene 120-12 Benzo(a)hijhoranthene 205-96 Benzo(b)fluoranthene 205-96 Benzo(b)fluoranthene 205-96 Benzo(b)fluoranthene 205-96 Benzo(b)fluoranthene 205-96 Benzo(b)fluoranthene 205-96 Benzo(a)hanthracene 191-24 Phenanthrene 191-24 Phenanthrene 191-20 Phenanthrene 1 | -7 6 -7 - -2 - -9 - -2 - -3 - -2 - -1 - -0 - -1 1 1 4 50 1 - -1 - | 0.014 5.6 16 990 0.79 1500 90 0.15 20 | 0.248 2.14 0.299 n/a 0.309 0.273 0.259 0.291 | 0.014 5.6 16 990 0.79 1500 |
|--|---|---|---|---|
| bis(2-Ethylhexyl)phthalate Butylbenzyl phthalate Caprolactam Dibenzofuran Dibenzofuran Dienthyl phthalate Din-butyl phthalate Din-butyl phthalate Din-octyl phthalate Hexachlorobenzene Hexachlorocyclopentadiene T7-47- Hexachlorocthane 67-72- Isophorone 78-59- Nitrobenzene n-Nitrosodimethylamine 62-75- n-Nitrosodimethylamine 62-16-40- n-Nitrosodin-propylamine 86-30- Pentachlorophenol Phenol Polycyclic Aromatic Hydrocarbons (PAHs)(ug/L) 1-Methylnaphthalene 2-Methylnaphthalene Acenaphthylene Acenaphtylene Acenaphtylene Acenaphtylene Acenaphtylene Acenaphtylen | -7 6 -7 - -2 - -9 - -2 - -3 - -2 - -1 - -0 - -1 1 1 4 50 1 - -1 - | 5.6 16 990 0.79 1500 90 0.15 20 | 2.14 0.299 n/a 0.309 0.273 0.259 | 5.6 16 990 0.79 |
| Butylbenzyl phthalate | -7 | 16 990 0.79 1500 90 0.15 20 | 0.299 n/a 0.309 0.273 0.259 | 16 990 0.79 |
| Caprolactam 105-60 | -2 - -9 - -2 - -3 - -2 - -1 - -0 - -1 1 1 4 50 1 - -1 - | 990 0.79 1500 90 0.15 20 | n/a 0.309 0.273 0.259 | 990 0.79 |
| Dibenzofuran 132-64 | -9 | 0.79 1500 90 0.15 20 | 0.309 0.273 0.259 | 0.79 |
| Diethyl phthalate | 2 - -3 - 2 - -1 - -0 - -1 1 1 4 50 1 - 1 - | 1500 90 0.15 20 | 0.273 0.259 | |
| Dimethyl phthalate | -3 | 90 0.15 20 | 0.259 | 1500 |
| Di-n-butyl phthalate 84-74-1 Dinitro-o-cresol (4,6-Dinitro-2-methylphenol) 534-52-1 Di-n-octyl phthalate 117-84-1 Hexachlorobenzene 118-74-1 Hexachlorocyclopentadiene 77-47-1 Hexachlorocyclopentadiene 67-72-1 Isophorone 78-59-1 Nitrobenzene 98-95-1 n-Nitrosodimethylamine 62-75-1 n-Nitrosodiphenylamine 86-30-1 Pentachlorophenol 87-86-1 Phenol 108-95 Polycyclic Aromatic Hydrocarbons (PAHs)(ug/L) 1-Methylnaphthalene 90-12-1 2-Methylnaphthalene 90-12-1 2-Methylnaphthalene 90-12-1 Acenaphthylene 83-32-1 Acenaphthylene 208-96 Anthracene 120-12 Benzo(a)anthracene 56-55-1 Benzo(b)fluoranthene 205-98-1 Benzo(b)fluoranthene 205-98-1 Benzo(g,h,i)perylene 191-24 Benzo(k)fluoranthene 53-70-1 Fluoranthene 53-70-1 Fluore | 2 - -1 - -0 - -1 1 1 4 50 1 - 1 - | 90 0.15 20 | (800-411(440) - 67, 578-43407) | |
| Dinitro-o-cresol (4,6-Dinitro-2-methylphenol) 534-52 Di-n-octyl phthalate 117-84 Hexachlorobenzene 118-74 Hexachlorocyclopentadiene 77-47-47-47-47-47-47-47-47-47-47-47-47-4 | -1 -0 -1 1 4 50 -1 -1 | 0.15 20 | 0.291 | H-H. |
| Dinitro-o-cresol (4,6-Dinitro-2-methylphenol) 534-52 Di-n-octyl phthalate 117-84 Hexachlorobenzene 118-74 Hexachlorocyclopentadiene 77-47-47-47-47-47-47-47-47-47-47-47-47-4 | -0 - -1 1 1 4 50 1 - 1 - | 20 | | 90 |
| Hexachlorobenzene | -1 1 4 50 -1 – | | 3.61 | 0.15 |
| Hexachlorobenzene | -1 1 4 50 -1 – | 0.0000 | 0.268 | 20 |
| Hexachlorocyclopentadiene | 4 50 1 – 1 – | 0.0098 | 0.280 | 0.0098 |
| Hexachloroethane 67-72. Isophorone 78-59. Nitrobenzene 98-95. n-Nitrosodimethylamine 621-64. n-Nitrosodiphenylamine 86-30. Pentachlorophenol 87-86. Phenol 108-95. Polycyclic Aromatic Hydrocarbons (PAHs)(ug/L) 1-Methylnaphthalene 90-12. 2-Methylnaphthalene 91-57. Acenaphthene 83-32. Acenaphthylene 208-96. Anthracene 120-12. Benzo(a)anthracene 56-55. Benzo(a)pyrene 50-32. Benzo(b)fluoranthene 205-99. Benzo(g,h,i)perylene 191-24. Benzo(k)fluoranthene 207-08. Chrysene 218-01. Dibenzo(a,h)anthracene 53-70. Fluoranthene 206-44. Fluorene 191-20. Indeno(1,2,3-c,d)pyrene 193-39. Naphthalene 91-20. Pyrene 129-00. Total Benzofluoranthenes (b,j,k) | 1 - | 0.041 | 1.08 | 0.041 |
| Isophorone | 1 | 0.33 | 0.300 | 0.33 |
| Nitrobenzene 98-95 n-Nitrosodimethylamine 62-75 n-Nitrosodiphenylamine 621-64 n-Nitrosodiphenylamine 86-30 Pentachlorophenol 87-86 Phenol 108-95 Polycyclic Aromatic Hydrocarbons (PAHs)(ug/L) 1-Methylnaphthalene 90-12 2-Methylnaphthalene 91-57 Acenaphthene 83-32 Acenaphthylene 208-96 Anthracene 120-12 Benzo(a)anthracene 56-55 Benzo(a)pyrene 50-32 Benzo(b)fluoranthene 205-99 Benzo(k)fluoranthene 207-08 Chrysene 218-01 Dibenzo(a,h)anthracene 53-70 Fluoranthene 53-70 Fluoranthene 91-20 Naphthalene 91-20 Pyene 129-00 Total Benzofluoranthenes (b,j,k) - Total HPAH - Total PAH - Polychlorinated Biphenyls (PCBs) (ug/L) | | 78 | 0.423 | 78 |
| n-Nitrosodimethylamine | 1975 | 0.14 | 0.253 | 0.14 |
| n-Nitrosodi-n-propylamine n-Nitrosodiphenylamine 86-30- Pentachlorophenol 87-86- Phenol 108-95 Polycyclic Aromatic Hydrocarbons (PAHs)(ug/L) 1-Methylnaphthalene 2-Methylnaphthalene 90-12- 2-Methylnaphthalene 83-32- Acenaphthene 83-32- Acenaphthylene 208-96 Anthracene 120-12 Benzo(a)anthracene 8enzo(a)pyrene 50-32- Benzo(b)fluoranthene 205-98 Benzo(b)fluoranthene 207-08 Chrysene Dibenzo(a,h)anthracene Fluoranthene 1206-44 Fluorene 1104-00 Naphthalene 91-20- Phenanthrene 91-20- Pyrene Total Benzofluoranthenes (b,j,k) Total HPAH Total LPAH Total PAH Total PAH Polychlorinated Biphenyls (PCBs) (ug/L) Aroclor 1016 12674-1 | (300) | 0.00011 | 1.33 | 0.00011 |
| n-Nitrosodiphenylamine 86-30 Pentachlorophenol 87-86 Phenol 108-95 Polycyclic Aromatic Hydrocarbons (PAHs)(ug/L) 1-Methylnaphthalene 90-12 2-Methylnaphthalene 91-57- Acenaphthene 83-32- Acenaphthylene 208-96 Anthracene 120-12 Benzo(a)anthracene 56-55- Benzo(a)pyrene 50-32- Benzo(b)fluoranthene 205-98 Benzo(g,h,i)perylene 191-24 Benzo(k)fluoranthene 207-08 Chrysene 218-01 Dibenzo(a,h)anthracene 53-70- Fluoranthene 206-44 Fluorene 193-39 Naphthalene 91-20- Phenanthrene 85-01- Pyrene 129-00 Total Benzofluoranthenes (b,j,k) - Total HPAH - Total PAH | | 0.011 | 0.269 | 0.011 |
| Pentachlorophenol 87-86-86 | | 12 | 0.299 | 12 |
| Phenol 108-95 Polycyclic Aromatic Hydrocarbons (PAHs)(ug/L) 90-12-12-12-12-12-12-12-12-12-12-12-12-12- | NO. | 0.041 | 1.89 | 0.041 |
| Polycyclic Aromatic Hydrocarbons (PAHs)(ug/L) 1-Methylnaphthalene 90-12-12-12-12-12-12-12-12-12-12-12-12-12- | 1.4g/ | 580 | 0.271 | 580 |
| 1-Methylnaphthalene 90-12- 2-Methylnaphthalene 91-57- Acenaphthene 83-32- Acenaphthylene 208-96- Anthracene 120-12- Benzo(a)anthracene 56-55- Benzo(b)fluoranthene 205-98- Benzo(b)fluoranthene 207-08- Benzo(k)fluoranthene 207-08- Chrysene 218-01- Dibenzo(a,h)anthracene 53-70- Fluoranthene 206-44 Fluorene 86-73- Indeno(1,2,3-c,d)pyrene 193-39- Naphthalene 91-20- Phenanthrene 85-01- Pyrene 129-00- Total Benzofluoranthenes (b,j,k) Total LPAH Total LPAH Total PAH Polychlorinated Biphenyls (PCBs) (ug/L) Aroclor 1016 12674-1 | -2 | 300 | 0.271 | 300 |
| 2-Methylnaphthalene 91-57-7-1 Acenaphthene 83-32-1 Acenaphthylene 208-96-1 Anthracene 120-12-1 Benzo(a)anthracene 56-55-1 Benzo(b)fluoranthene 205-98-1 Benzo(b)fluoranthene 207-08-1 Benzo(k)fluoranthene 207-08-1 Chrysene 218-01-1 Dibenzo(a,h)anthracene 53-70-1 Fluoranthene 206-44-1 Fluorene 86-73-1 Indeno(1,2,3-c,d)pyrene 193-39-1 Naphthalene 91-20-1 Pyrene 129-00-1 Total Benzofluoranthenes (b,j,k) | n I | 1.1 | 0.0100 | 1.1 |
| Acenaphthene 83-32-32-32-32-32-32-32-32-32-32-32-32-32 | | 3.6 | 0.0100 | 3.6 |
| Acenaphthylene 208-96 Anthracene 120-12 Benzo(a)anthracene 56-55 Benzo(b)fluoranthene 205-99 Benzo(g,h,i)perylene 191-24 Benzo(k)fluoranthene 207-08 Chrysene 218-01 Dibenzo(a,h)anthracene 53-70 Fluoranthene 206-44 Fluorene 86-73 Indeno(1,2,3-c,d)pyrene 193-39 Naphthalene 91-20 Phenanthrene 85-01 Pyrene 129-00 Total Benzofluoranthenes (b,j,k) Total LPAH Total PAH Polychlorinated Biphenyls (PCBs) (ug/L) Aroclor 1016 12674-1 | | 53 | 0.0100 | 53 |
| Anthracene 120-12 Benzo(a)anthracene 56-55 Benzo(a)pyrene 50-32 Benzo(b)fluoranthene 205-99 Benzo(k)fluoranthene 191-24 Benzo(k)fluoranthene 207-08 Chrysene 218-01 Dibenzo(a,h)anthracene 53-70- Fluoranthene 206-44 Fluorene 86-73- Indeno(1,2,3-c,d)pyrene 193-39 Naphthalene 91-20- Phenanthrene 85-01- Pyrene 129-00 Total Benzofluoranthenes (b,j,k) - Total HPAH - Total PAH - Polychlorinated Biphenyls (PCBs) (ug/L) Aroclor 1016 12674-1 | | 53 | 0.0100 | |
| Benzo(a)anthracene 56-55-56-56-56-56-56-56-56-56-56-56-56-5 | | | | 400 |
| Benzo(a)pyrene 50-32-32-32-32-32-32-32-32-32-32-32-32-32- | | 180 | 0.0100 | 180 |
| Benzo(b)fluoranthene 205-99 Benzo(g,h,i)perylene 191-24 Benzo(k)fluoranthene 207-08 Chrysene 218-01 Dibenzo(a,h)anthracene 53-70- Fluoranthene 206-44 Fluorene 86-73- Indeno(1,2,3-c,d)pyrene 193-39 Naphthalene 91-20- Phenanthrene 85-01- Pyrene 129-00 Total Benzofluoranthenes (b,j,k) - Total LPAH - Total PAH - Polychlorinated Biphenyls (PCBs) (ug/L) Aroclor 1016 12674-1 | | 0.012 | 0.0100 | 0.012 |
| Benzo(g,h,i)perylene 191-24 Benzo(k)fluoranthene 207-08 Chrysene 218-01 Dibenzo(a,h)anthracene 53-70-1 Fluoranthene 206-44 Fluorene 86-73-1 Indeno(1,2,3-c,d)pyrene 193-39-39-39-39-39-39-39-39-39-39-39-39-3 | | 0.0034 | 0.0100 | 0.0034 |
| Benzo(k)fluoranthene 207-08 Chrysene 218-01 Dibenzo(a,h)anthracene 53-70-1 Fluoranthene 206-44 Fluorene 86-73-1 Indeno(1,2,3-c,d)pyrene 193-39 Naphthalene 91-20-1 Phenanthrene 85-01-1 Pyrene 129-00 Total Benzofluoranthenes (b,j,k) - Total HPAH - Total LPAH - Total PAH - Polychlorinated Biphenyls (PCBs) (ug/L) Aroclor 1016 12674-1 | | 0.034 | 0.0100 | 0.034 |
| Chrysene 218-01 Dibenzo(a,h)anthracene 53-70- Fluoranthene 206-44 Fluorene 86-73- Indeno(1,2,3-c,d)pyrene 193-39- Naphthalene 91-20- Phenanthrene 85-01- Pyrene 129-00- Total Benzofluoranthenes (b,j,k) Total HPAH Total LPAH Total PAH Polychlorinated Biphenyls (PCBs) (ug/L) Aroclor 1016 12674-1 | | | 0.0100 | |
| Dibenzo(a,h)anthracene 53-70- Fluoranthene 206-44 Fluorene 86-73- Indeno(1,2,3-c,d)pyrene 193-39- Naphthalene 91-20- Phenanthrene 85-01- Pyrene 129-00- Total Benzofluoranthenes (b,j,k) - Total HPAH - Total LPAH - Total PAH - Polychlorinated Biphenyls (PCBs) (ug/L) 12674-1 | | 0.34 | 0.0100 | 0.34 |
| Fluoranthene 206-44 Fluorene 86-73-1 Indeno(1,2,3-c,d)pyrene 193-39 Naphthalene 91-20-1 Phenanthrene 85-01-1 Pyrene 129-00-1 Total Benzofluoranthenes (b,j,k) | | 3.4 | 0.0100 | 3.4 |
| Fluorene 86-73-1 Indeno(1,2,3-c,d)pyrene 193-39 Naphthalene 91-20-1 Phenanthrene 85-01-1 Pyrene 129-00-1 Total Benzofluoranthenes (b,j,k) | | 0.0034 | 0.0100 | 0.0034 |
| Indeno(1,2,3-c,d)pyrene 193-39 Naphthalene 91-20- Phenanthrene 85-01- Pyrene 129-00 Total Benzofluoranthenes (b,j,k) Total HPAH Total LPAH Total PAH Polychlorinated Biphenyls (PCBs) (ug/L) Aroclor 1016 12674-1 | | 80 | 0.0100 | 80 |
| Naphthalene 91-20- Phenanthrene 85-01- Pyrene 129-00 Total Benzofluoranthenes (b,j,k) Total HPAH Total LPAH Total PAH Polychlorinated Biphenyls (PCBs) (ug/L) Aroclor 1016 12674-1 | | 29 | 0.0100 | 29 |
| Phenanthrene 85-01- Pyrene 129-00 Total Benzofluoranthenes (b,j,k) Total HPAH Total LPAH Total PAH Polychlorinated Biphenyls (PCBs) (ug/L) Aroclor 1016 12674-1 | | 0.034 | 0.0100 | 0.034 |
| Pyrene 129-00 Total Benzofluoranthenes (b,j,k) Total HPAH Total LPAH Total PAH Polychlorinated Biphenyls (PCBs) (ug/L) Aroclor 1016 12674-1 | 3 – | 0.17 | 0.0100 | 0.17 |
| Total Benzofluoranthenes (b,j,k) — Total HPAH — Total LPAH — Total PAH — Polychlorinated Biphenyls (PCBs) (ug/L) Aroclor 1016 12674-1 | 8 - | | 0.0100 | |
| Total HPAH — Total LPAH — Total PAH — Polychlorinated Biphenyls (PCBs) (ug/L) Aroclor 1016 12674-1 | -0 - | 12 | 0.0100 | 12 |
| Total LPAH Total PAH Polychlorinated Biphenyls (PCBs) (ug/L) Aroclor 1016 12674-1 | - | | | , - |
| Total PAH – Polychlorinated Biphenyls (PCBs) (ug/L) Aroclor 1016 12674-1 | | <u> </u> | <u> </u> | |
| Polychlorinated Biphenyls (PCBs) (ug/L) Aroclor 1016 12674-1 | _ | | | 12 <u>20</u> 0 |
| Aroclor 1016 12674-1 | _ | | - | - |
| | - | • | | |
| Aroclor 1221 11104-2 | - | 0.14 | 1.0 | 0.14 |
| | 1-2 _ | 0.0047 | 1.0 | 0.0047 |
| Aroclor 1232 11141-1 | | 0.0047 | 1.0 | 0.0047 |
| Aroclor 1242 53469-2 | 8-2 | 0.0078 | 1.0 | 0.0078 |
| Aroclor 1248 12672-2 | 8-2 6-5 | | 1.0 | 0.0078 |
| Aroclor 1254 11097-6 | 8-2 — 6-5 — 1-9 — | | 1.0 | 0.0078 |
| Aroclor 1260 11096-8 | 8-2 6-5 1-9 9-6 | 0.0078 | 1.0 | 0.0078 |
| Total PCB Aroclors 1336-36 | 8-2 — 6-5 — 1-9 — 9-6 — 9-1 — | | | 0.044 |

Notes:

Compounds frequently associated with MGP-operations.

'-- indicates not available

CAS = Chemical Abstract Services

EPA = U. S. Environmental Protection Agency

L = liter

MCL = maximum contaminant level

mg = miligram

MGP = manufactured gas plant

ng = nanogram

PRG = preliminary remediation goal

RSL = regional screening level

ug = microgram

References:

EPA, 2016. EPA Regional Screening Levels. May 2016. Available from: https://www.epa.gov/risk/regional-screening-levels-rsls-generic-tables-may-2016

Table 3-6 - Development of Initial PRGs for Sediment Bremerton Gas Works Superfund Site Bremerton, Washington

| | | SMS Marine Sediment | SMS Marine Cleanup | EDA Darian E DCDA | FDA Darian 2 DTAC | | | |
|--|--|--|---|--|--|----------------------------|--------------------------------|--|
| Analyte | | Cleanup Objective (SCO ¹ /LAET ²) | Screening Level (CSL ¹ /2LAET ²) | EPA Region 5 RCRA Sediment Ecological Screening Levels | EPA Region 3 BTAG Marine Sediment Screening Benchmarks | Effects Range-Low (ERL) | Effects Range- Median (ERM) | Initial PRGs used for Data |
| | CAS Number | DOE, 2013 | DOE, 2013 | EPA, 2003 | EPA, 2006 | Long et al., 1995 | Long et al., 1995 | Screening ³ |
| Alkane Isomers (ug/kg) n-Hexane (C6) | 110-54-3 | 122 | | | 39.60 | | | 39.6 |
| Conventionals (mg/kg) | 110-54-5 | | | | 39.00 | | | 39.0 |
| Cyanide, WAD | 57-12-5 | | | | 0.1 | | | 0.1 |
| Cyanide, total | 57-12-5 | | | 0.0001 | | | | 0.0001 |
| Sulfide Metals (mg/kg) | 18496-25-8 | | | | 130 | | | 130 |
| Antimony | 7440-36-0 | | | | 2 | | | 2 |
| Arsenic | 7440-38-2 | 57 | 93 | 9.79 | 7.24 | 8.2 | 70 | 57 |
| Beryllium | 7440-41-7 | | | | | | | |
| Cadmium Chromium | 7440-43-9 7440-47-3 | 5.1 260 | 6.7 270 | 0.99 43.4 | 0.68 52.3 | 1.2 81 | 9.6 370 | 5.1 260 |
| Chromium III | 16065-83-1 | | | | | | | |
| Chromium VI | 18540-29-9 | | | | | | | 1 |
| Copper | 7440-50-8 | 390 | 390 | 31.6 | 18.7 | 34 | 270 | 390 |
| Lead Mercury | 7439-92-1 7439-97-6 | 450 0.41 | 530 0.59 | 35.8 0.174 | 30.2 0.13 | 46.7 0.15 | 218 0.71 | 450 0.41 |
| Nickel | 7439-97-6 | U.41 | 0.59 | 22.7 | 15.9 | 20.9 | 51.6 | 20.9 |
| Selenium | 7782-49-2 | | | | 2 | | | 2 |
| Silver | 7440-22-4 | 6.1 | 6.1 | 0.5 | 0.73 | 1 | 3.7 | 6.1 |
| Thallium | 7440-28-0 | | | | | | | |
| Zinc Metals, Organic (ug/kg) | 7440-66-6 | 410 | 960 | 121 | 124 | 150 | 410 | 410 |
| Tributyltin | 688-73-3 | | | | - | | | |
| Polycyclic Aromatic Hydrocarbons (PAHs) (ug/kg) | | | 1 | 1 | | | | |
| 1-Methylnaphthalene | 90-12-0 | | | | | | | - 1 |
| 2-Methylnaphthalene | 91-57-6 | 670 | 670 | 20.2 | 20.2 | 70 | 670 | 670 |
| Acenaphthylene | 83-32-9 | 500 | 500 1300 | 6.71 5.87 | 6.71 5.87 | 16 | 500 | 500 1300 |
| Acenaphthylene Anthracene | 208-96-8 120-12-7 | 1300 960 | 1300 960 | 5.87 57.2 | 5.87 46.9 | 44 85.3 | 640 1100 | 1300 960 |
| Benzo(a)anthracene | 56-55-3 | 1300 | 1600 | 108 | 74.8 | 261 | 1600 | 1300 |
| Benzo(a)pyrene | 50-32-8 | 1600 | 1600 | 150 | 88.8 | 430 | 1600 | 1600 |
| Benzo(b)fluoranthene | 205-99-2 | | | 10400 | | | | 10400 |
| Benzo(b,k)fluoranthene | 101 24 2 | | 720 | 170 | 27.2 | 13 13 . | ' | 27.2 |
| Benzo(g,h,i)perylene Benzo(j)fluoranthene | 191-24-2 205-82-3 | 670 | 720 | 170 | 170 | | | 670 |
| Benzo(k)fluoranthene | 207-08-9 | | | 240 | 240 | | | 240 |
| Chrysene | 218-01-9 | 1400 | 2800 | 166 | 108 | 384 | 2800 | 1400 |
| Dibenzo(a,h)anthracene | 53-70-3 | 230 | 230 | 33 | 6.22 | 63.4 | 260 | 230 |
| Fluoranthene | 206-44-0 | 1700 | 2500 | 423 | 113 | 600 | 5100 | 1700 |
| Fluorene Indeno(1,2,3-c,d)pyrene | 86-73-7 193-39-5 | 540 600 | 540 690 | 77.4 200 | 21.2 17 | 19 | 540 | 540 600 |
| Naphthalene | 91-20-3 | 2100 | 2100 | 176 | 34.6 | 160 | 2100 | 2100 |
| Phenanthrene | 85-01-8 | 1500 | 1500 | 204 | 86.7 | 240 | 1500 | 1500 |
| Pyrene | 129-00-0 | 2600 | 3300 | 195 | 153 | 665 | 2600 | 2600 |
| Total Benzofluoranthenes (b,j,k) Total HPAH | | 3200 | 3600 | | 655 | 1700 | 9600 | 3200 |
| Total LPAH | | 12000 5200 | 17000 5200 | | 312 | 552 | 3160 | 12000 5200 |
| Total PAH | | | | | 2900 | 4022 | 44792 | 4022 |
| Polycyclic Aromatic Hydrocarbons (PAHs) (mg/kg-OC) | | | | | | | | |
| 2-Methylnaphthalene | 91-57-6 | 38 | 64 | | | | | 38 |
| Acenaphthene Acenaphthylene | 83-32-9 208-96-8 | 16 66 | 57 66 | | | | | 16 66 |
| Anthracene | 120-12-7 | 220 | 1200 | | | | | 220 |
| Benzo(a)anthracene | 56-55-3 | 110 | 270 | | | | | 110 |
| Benzo(a)pyrene | 50-32-8 | 99 | 210 | | 73 | - | - | 99 |
| Benzo(g,h,i)perylene | 191-24-2 | 31 | 78 | | .=- | | | 31 |
| Chrysene Dibenzo(a,h)anthracene | 218-01-9 53-70-3 | 110 12 | 460 33 | | | | | 110 12 |
| Fluoranthene | 206-44-0 | 160 | 1200 | | | | - | 160 |
| Fluorene | 86-73-7 | 23 | 79 | | = | | | 23 |
| Indeno(1,2,3-c,d)pyrene | 193-39-5 | 34 | 88 | | | | | 34 |
| Naphthalene Phenanthrene | 91-20-3 85-01-8 | 99 | 170 480 | | | | | 99 100 |
| Pyrene | 129-00-0 | 1000 | 1400 | | | | | 1000 |
| Total Benzofluoranthenes (b,j,k) | - | 230 | 450 | | | | | 230 |
| Total HPAH | | 960 | 5300 | | | - | | 960 |
| Total LPAH Rehyelleringted Riphonyle (RCRs) (ug/kg) | | 370 | 780 | - ni- | | | <u></u> | 370 |
| Polychlorinated Biphenyls (PCBs) (ug/kg) Aroclor 1016 | 12674-11-2 | | | | | - | | |
| Aroclor 1221 | 11104-28-2 | | | | | | | |
| Aroclor 1232 | 11141-16-5 | 1 | | | 74 | == | | - |
| Aroclor 1242 | 53469-21-9 | | | | | , | | - |
| Aroclor 1248 Aroclor 1254 | 12672-29-6 11097-69-1 | | - | | 63.3 | | | 62.2 |
| Aroclor 1254 Aroclor 1260 | 11027-03-1 | | | | 63.3 | | | 63.3 |
| Aroclor 1262 | A CONTRACTOR OF THE PARTY OF TH | | | | | | | |
| A 00101 1202 | 11096-82-5 37324-23-5 | | | | | Sv. | | |
| Aroclor 1268 | 11096-82-5 | | | | | | ļ | |
| Aroclor 1268 Total PCB Aroclors | 11096-82-5 37324-23-5 | | | 59.8 | 40 | 22.7 | 180 | 130 |
| Aroclor 1268 Total PCB Aroclors Polychlorinated Biphenyls (PCBs) (mg/kg-OC) | 11096-82-5 37324-23-5 11100-14-4 | 130 | 1000 | 59.8 | 40 | 22.7 | 180 | 130 |
| Aroclor 1268 Total PCB Aroclors Polychlorinated Biphenyls (PCBs) (mg/kg-OC) Total PCB Aroclors | 11096-82-5 37324-23-5 11100-14-4 | | | | | | - | |
| Aroclor 1268 Total PCB Aroclors Polychlorinated Biphenyls (PCBs) (mg/kg-OC) | 11096-82-5 37324-23-5 11100-14-4 | 130 | 1000 | 59.8 | 40 | 22.7 | 180 | 130 |
| Aroclor 1268 Total PCB Aroclors Polychlorinated Biphenyls (PCBs) (mg/kg-OC) Total PCB Aroclors Semivolatile Organic Compounds (SVOCs)(ug/kg) 1,2,4,5-Tetrachlorobenzene 1,2,4-Trichlorobenzene | 11096-82-5 37324-23-5 11100-14-4 95-94-3 120-82-1 | 130 12 31 | 1000 65 51 | 59.8 1252 5062 | 47000 473 | 22.7 | 180 | 130 12 47000 31 |
| Aroclor 1268 Total PCB Aroclors Polychlorinated Biphenyls (PCBs) (mg/kg-OC) Total PCB Aroclors Semivolatile Organic Compounds (SVOCs)(ug/kg) 1,2,4,5-Tetrachlorobenzene 1,2,4-Trichlorobenzene 1,2-Dichlorobenzene | 11096-82-5 37324-23-5 11100-14-4 95-94-3 120-82-1 95-50-1 | 130 12 31 35 | 1000 65 51 50 | 59.8 1252 5062 294 | 40 47000 473 989 | | | 130 12 47000 31 35 |
| Aroclor 1268 Total PCB Aroclors Polychlorinated Biphenyls (PCBs) (mg/kg-OC) Total PCB Aroclors Semivolatile Organic Compounds (SVOCs)(ug/kg) 1,2,4,5-Tetrachlorobenzene 1,2,4-Trichlorobenzene 1,2-Dichlorobenzene 1,3-Dichlorobenzene | 11096-82-5 37324-23-5 11100-14-4 95-94-3 120-82-1 95-50-1 541-73-1 | 130 12 31 35 | 1000 65 51 50 | 59.8 1252 5062 294 1315 | 47000 473 989 842 | | | 130 12 47000 31 35 842 |
| Aroclor 1268 Total PCB Aroclors Polychlorinated Biphenyls (PCBs) (mg/kg-OC) Total PCB Aroclors Semivolatile Organic Compounds (SVOCs)(ug/kg) 1,2,4,5-Tetrachlorobenzene 1,2,4-Trichlorobenzene 1,2-Dichlorobenzene 1,3-Dichlorobenzene 1,4-Dichlorobenzene | 11096-82-5 37324-23-5 11100-14-4 95-94-3 120-82-1 95-50-1 541-73-1 106-46-7 | 130 12 31 35 | 1000 65 51 50 | 59.8 1252 5062 294 | 40 47000 473 989 | | | 130 12 47000 31 35 842 110 |
| Aroclor 1268 Total PCB Aroclors Polychlorinated Biphenyls (PCBs) (mg/kg-OC) Total PCB Aroclors Semivolatile Organic Compounds (SVOCs)(ug/kg) 1,2,4,5-Tetrachlorobenzene 1,2,4-Trichlorobenzene 1,2-Dichlorobenzene 1,3-Dichlorobenzene | 11096-82-5 37324-23-5 11100-14-4 95-94-3 120-82-1 95-50-1 541-73-1 | 130 12 31 35 110 | 1000 65 51 50 110 | 59.8 1252 5062 294 1315 318 | 40 47000 473 989 842 460 | | | 130 12 47000 31 35 842 |
| Aroclor 1268 Total PCB Aroclors Polychlorinated Biphenyls (PCBs) (mg/kg-OC) Total PCB Aroclors Semivolatile Organic Compounds (SVOCs)(ug/kg) 1,2,4,5-Tetrachlorobenzene 1,2,4-Trichlorobenzene 1,2-Dichlorobenzene 1,3-Dichlorobenzene 1,4-Dichlorobenzene 2,2'-Oxybis (1-chloropropane) | 11096-82-5 37324-23-5 11100-14-4 95-94-3 120-82-1 95-50-1 541-73-1 106-46-7 108-60-1 | 130 12 31 35 110 | 1000 65 51 50 110 | 59.8 1252 5062 294 1315 318 | 47000 473 989 842 460 | | | 130 12 47000 31 35 842 110 |

Table 3-6 - Development of Initial PRGs for Sediment

Bremerton Gas Works Superfund Site Bremerton, Washington

SMS Marine Sediment **SMS Marine Cleanup EPA Region 5 RCRA EPA Region 3 BTAG Screening Level Cleanup Objective Effects Range-Low** Initial PRGs used Sediment Ecological **Marine Sediment Effects Range-**(SCO¹/LAET²) (CSL¹/2LAET²) **Analyte Screening Levels Screening Benchmarks** (ERL) Median (ERM) for Data DOE, 2013 DOE, 2013 EPA, 2003 **EPA, 2006** Long et al., 1995 Long et al., 1995 **CAS Number** Screening³ 120-83-2 2,4-Dichlorophenol 81.7 117 117 2,4-Dimethylphenol 105-67-9 29 29 304 29 -----29 51-28-5 4.4-Dinitrophenol 6.21 __ _ --6.21 ,4-Dinitrotoluene 121-14-2 14.4 41.6 41.6 39.8 ,6-Dinitrotoluene 606-20-2 ---------39.8 -Chloronaphthalene 91-58-7 417 417 -Chlorophenol 95-57-8 31.9 344 344 . .---. !-Methylphenol (o-Cresol) 95-48-7 63 63 55.4 --------63 88-74-4 -Nitroaniline -----------------Nitrophenol 88-75-5 -----3,3'-Dichlorobenzidine 91-94-1 127 2060 2060 3-Methylphenol & 4-Methylphenol (m&p-Cresol) 1319-77-3 -----52.4 3-Methylphenol (m-Cresol) 108-39-4 52.4 99-09-2 -Nitroaniline --------.... -I-Bromophenyl-phenyl ether 101-55-3 1550 1230 1230 -----Chloro-3-methylphenol 59-50-7 388 388 __ 146 4-Chloroaniline 106-47-8 ___ ___ __ __ __ 146 4-Methylphenol (p-Cresol) 106-44-5 670 670 20.2 670 670 4-Nitroaniline 100-01-6 ---------------100-02-7 13.3 4-Nitrophenol 13.3 98-86-2 Acetophenone -------------62-53-3 0.31 Aniline --0.31 --1912-24-9 6.62 Atrazine ---6.62 Benzaldehyde 100-52-7 ----------__ --Benzidine 92-87-5 Benzoic acid 65-85-0 650 650 650 650 -------100-51-6 57 73 1.04 Benzyl alcohol --57 1220 Biphenyl (1,1'-Biphenyl) 92-52-4 1220 -----------bis(2-Chloroethoxy)methane 111-91-1 -----------111-44-4 bis(2-Chloroethyl)ether 3520 3520 bis(2-Ethylhexyl)phthalate 117-81-7 1300 3100 182 182 1300 ----85-68-7 1970 Butylbenzyl phthalate 63 900 16800 63 105-60-2 Caprolactam -------------132-64-9 540 540 449 7300 Dibenzofuran 540 Diethyl phthalate 84-66-2 200 1200 295 218 200 131-11-3 160 Dimethyl phthalate 71 ------__ 71 84-74-2 1400 5100 1114 Di-n-butyl phthalate 1160 1400 Dinitro-o-cresol (4,6-Dinitro-2-methylphenol) 534-52-1 104 --104 -----------Di-n-octyl phthalate 117-84-0 6200 6200 40600 ----6200 118-74-1 70 20 Hexachlorobenzene 22 20 22 77-47-4 --901 139 139 Hexachlorocyclopentadiene ----67-72-1 584 804 **Hexachloroethane** 804 432 sophorone 78-59-1 -------------432 98-95-3 145 145 Nitrobenzene 62-75-9 n-Nitrosodimethylamine .----------. -n-Nitrosodi-n-propylamine 621-64-7 ----n-Nitrosodiphenylamine 86-30-6 28 40 422000 28 --360 690 7970 87-86-5 23000 Pentachlorophenol ----360 108-95-2 420 1200 49.1 420 Phenol 420 |Semivolatile Organic Compounds (SVOCs) (mg/kg-OC) 1,2,4-Trichlorobenzene 120-82-1 0.81 1.8 0.81 1,2-Dichlorobenzene 95-50-1 2.3 2.3 2.3 ---------1,4-Dichlorobenzene 106-46-7 3.1 9 ----3.1 bis(2-Ethylhexyl)phthalate 117-81-7 47 78 47 4.9 64 Butylbenzyl phthalate 85-68-7 -----------4.9 Dibenzofuran 132-64-9 15 58 15 ---Diethyl phthalate 61 110 84-66-2 61 ---------131-11-3 53 53 Dimethyl phthalate 53 ----Di-n-butyl phthalate 84-74-2 220 1700 220 __ ----117-84-0 58 4500 Di-n-octyl phthalate ---__ 58 __ 118-74-1 0.38 2.3 Hexachlorobenzene ---0.38 11 11 n-Nitrosodiphenylamine 86-30-6 -------11 --Volatile Organic Compounds (VOCs) (ug/kg) 1,1,1,2-Tetrachloroethane 630-20-6 ----1,1,1-Trichloroethane 71-55-6 213 856 856 .----850 202 1,1,2,2-Tetrachloroethane 79-34-5 202 518 570 79-00-5 1,1,2-Trichloroethane ---------570 1,1,2-Trichlorotrifluoroethane (Freon 113) 76-13-1 1,1-Dichloroethane 75-34-3 0.575 0.575 --------------75-35-4 2780 1,1-Dichloroethene __ 19.4 --2780 --1,2,3-Trichlorobenzene 87-61-6 858 ------858 1,2,3-Trichloropropane 96-18-4 ------------95-63-6 1,2,4-Trimethylbenzene __ 1,2-Dibromo-3-chloropropane 96-12-8 .--------------.... ---107-06-2 260 1,2-Dichloroethane -------260 ---1,2-Dichloroethene, cis-156-59-2 -----------------1,2-Dichloroethene, trans-156-60-5 654 1050 --1050 78-87-5 333 L,2-Dichloropropane 333 108-67-8 1,3,5-Trimethylbenzene (Mesitylene) __ -------__ --142-28-9 1,3-Dichloropropane ------1,3-Dichloropropene, cis-10061-01-5 y.----- --------------__ 1,3-Dichloropropene, trans-10061-02-6 __ --------110-57-6 .,4-Dichloro-2-butene, trans------------1,4-Dioxane 123-91-1 119 119 --------Butanone (MEK) 78-93-3 42.4 42.4 -Chlorotoluene 95-49-8 -------------58.2 !-Hexanone (Methyl butyl ketone) 591-78-6 58.2 1-Chlorotoluene 106-43-4 -----------------99-87-6 4-Isopropyltoluene (4-Cymene) __ __ __ ___ 67-64-1 9.9 9.9 Acetone --107-02-8 0.00152 Acrolein ---------0.00152 Acrylonitrile 107-13-1 1.2 1.2

Table 3-6 - Development of Initial PRGs for Sediment

Bremerton Gas Works Superfund Site

Bremerton, Washington

| Analyte | CAS Number | SMS Marine Sediment Cleanup Objective (SCO ¹ /LAET ²) DOE, 2013 | SMS Marine Cleanup Screening Level (CSL ¹ /2LAET ²) DOE, 2013 | EPA Region 5 RCRA Sediment Ecological Screening Levels EPA, 2003 | EPA Region 3 BTAG Marine Sediment Screening Benchmarks EPA, 2006 | Effects Range-Low (ERL) Long et al., 1995 | Effects Range- Median (ERM) Long et al., 1995 | Initial PRGs used for Data Screening ³ |
|---|------------|---|---|---|---|---|---|---|
| Benzene | 71-43-2 | | | 142 | 137 | | : | 137 |
| Bromobenzene | 108-86-1 | : | | | | | | |
| Bromochloromethane | 74-97-5 | | | | | ##A | 144 | |
| Bromodichloromethane | 75-27-4 | 1 | : | | | | | |
| Bromoform (Tribromomethane) | 75-25-2 | | | 492 | 1310 | | | 1310 |
| Bromomethane (Methyl bromide) | 74-83-9 | | - | 1.37 | | | i_u | 1.37 |
| Carbon disulfide | 75-15-0 | | | 23.9 | 0.851 | | | 0.851 |
| Carbon tetrachloride (Tetrachloromethane) | 56-23-5 | F | for-them (in-the) | 1450 | 7240 | | | 7240 |
| Chlorobenzene | 108-90-7 | : | | 291 | 162 | | 1000 | 162 |
| Chloroethane | 75-00-3 | | | | -u | ₩-1 | 1-1 | |
| Chloroform | 67-66-3 | 1 | | 121 | | | 1 | 121 |
| Chloromethane | 74-87-3 | | E# | == | | #= | | |
| Cyclohexane | 110-82-7 | | | | | | | |
| Dibromochloromethane | 124-48-1 | | | | | | | |
| Dibromomethane | 74-95-3 | | == | | | | | |
| Dichlorodifluoromethane | 75-71-8 | | | | | | | |
| Dichloromethane (Methylene chloride) | 75-09-2 | | | 159 | | | | 159 |
| Ethylbenzene | 100-41-4 | | | 175 | 305 | | | 305 |
| Ethylene dibromide (1,2-Dibromoethane) | 106-93-4 | | | | | | ; | |
| Hexachlorobutadiene (Hexachloro-1,3-butadiene) | 87-68-3 | 11 | 120 | 26.5 | | | | 11 |
| Isopropylbenzene (Cumene) | 98-82-8 |) Ho | (HII | | 86 | | 3-8 | 86 |
| Methyl acetate | 79-20-9 | | | | | | | |
| Methyl iodide (lodomethane) | 74-88-4 | | - | | | | | |
| Methyl isobutyl ketone (4-Methyl-2-pentanone or (MIBK)) | 108-10-1 | | | 25.1 | | | | 25.1 |
| Methyl tert-butyl ether (MTBE) | 1634-04-4 | | | | | | | |
| n-Butylbenzene | 104-51-8 | | | | | | | |
| n-Propylbenzene | 103-65-1 | | | | | | | |
| o-Xylene | 95-47-6 | | | | | | | |
| sec-Butylbenzene | 135-98-8 | | | | | | | |
| Styrene | 100-42-5 | | | 254 | 7070 | | 1 | 7070 |
| tert-Butylbenzene | 98-06-6 | | | | - 1770A (FO) | | | |
| Tetrachloroethene (PCE) | 127-18-4 | | | 990 | 190 | | | 190 |
| Toluene | 108-88-3 | | | 1220 | 1090 | | | 1090 |
| Total xylene (reported, not calculated) | 1330-20-7 | | - | 433 | | | | 433 |
| Total Xylene | | | | 433 | | | | 433 |
| Trichloroethene (TCE) | 79-01-6 | | | 112 | 8950 | | | 8950 |
| Trichlorofluoromethane (Fluorotrichloromethane) | 75-69-4 | | | | | | | |
| Vinyl acetate | 108-05-4 | | | 13 | | | | 13 |
| Vinyl chloride | 75-01-4 | | | 202 | | - | H-1 | 202 |
| Volatile Organic Compounds (VOCs) (mg/kg-OC) | | | | -9- | | | | |
| Hexachlorobutadiene (Hexachloro-1,3-butadiene) | 87-68-3 | 3.9 | 6.2 | | | | 144 | 3.9 |
| Notes: | | 100 (100) | | | | | | |

Notes:

Compounds frequently associated with MGP-operations

- '-- indicates not available
- 1 = This criteria will be used when total organic carbon (TOC) is between 0.5% to 5%.
- 2 = This criteria will be used when total organic carbon (TOC) is less than 0.5% or greater than 5%.
- 3 = Site-specific fish and shellfish consumption based PRGs have not yet been developed. The fish and shellfish based PRGs will be developed in consultation with EPA and the Suquamish Tribe as part of this RI/FS Work Plan implementation.

2LAET = Second Lowest Apparent Effects Threshold

BTAG = Biological Technical Assistance Group

CAS = Chemical Abstract Services

CSL = Cleanup Screening Level DOE = Washington Department of Ecology

EPA = United States Environmental Protection Agency

kg = kilogram LAET = Lowest Apparent Effects Threshold

mg = miligram

MGP = Manufactured Gas Plant

ng = nanogram OC = organic carbon

PRG = preliminary remediation goal

RCRA = Resource Conservation and Recovery Act

SCO = Sediment Cleanup Objective

SMS = Sediment Management Standards

ug = microgram

References:

Ecology, 2013. Sediment Management Standards, Chapter 173-204 WAC: Final Rule February 22, 2013. September 1, 2013.

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Table 3-7 - Development of Initial PRGs for Surface Water

| | | | AR | ARs | | Other Screening B | enchmark Sources | |
|--|------------|---|---|--|---|---|---|----------------------------|
| | | National Recommended Water Quality Criteria - Aquatic Life Criteria - Saltwater CCC (chronic) ¹ | National Recommended Water Quality Criteria - Aquatic Life Criteria - Saltwater CMC (acute) ¹ | National Recommended Water Quality Criteria - Human Health for the Consumption of Organisms | EPA 2016 CWA- Effective Human Health Criteria Applicable to Washington (Organism Only) | EPA Region 3 BTAG Marine Water Screening Benchmarks | EPA Region 5 RCRA - Ecological Screening Levels - Water | Initial PRGs used for Data |
| Analyte | CAS Number | EPA, 2013 | EPA, 2013 | EPA, 2013 | EPA/Ecology,2016 | EPA, 2006 | EPA, 2003 | Screening |
| Alkane Isomers (ug/L) | | | _ | | | | | |
| n-Hexane (C6) | 110-54-3 | | | - | | 0.58 | H | 0.58 |
| Conventionals (mg/L) | | | _ | | | | | |
| Cyanide, free | 57-12-5 | 0.001 | 0.001 | | | 0.001 | | 0.001 |
| Cyanide, total | 57-12-5 | | == | 0.14 | 0.10 | : | 0.0052 | 0.10 |
| Sulfide | 18496-25-8 | = | | \ | == | = | | |
| Metals (ug/L) | | | | | | | | |
| Antimony | 7440-36-0 | == | | 640 | 90 | 500 | 80 | 90 |
| Arsenic | 7440-38-2 | 36 | 69 | 0.14 | 0.14 | 12.5 (a) | 148 | 0.14 |
| Beryllium | 7440-41-7 | | | | | 0.66 | 3.6 | 0.66 |
| Cadmium | 7440-43-9 | 8.8 | 40 | | | 0.12 (a) | 0.15 | 8.8 |
| Chromium | 7440-47-3 | | - | | | 57.5 | 42 | 42 |
| Chromium III | 16065-83-1 | = | | i=- | | 56 (a) | | 56 |
| Chromium VI | 18540-29-9 | 50 | 1100 | H | 99 | 1.5 (a) | H | 50 |
| Copper | 7440-50-8 | 3.1 | 4.8 | H- | | 3.1 | 1.58 | 3.1 |
| Lead | 7439-92-1 | 8.1 | 210 | | 22 | 8.1 | 1.17 | 8.1 |
| Mercury | 7439-97-6 | 0.94 | 1.8 | F4- | | 0.016 (a) | 0.0013 | 0.94 |
| Nickel | 7440-02-0 | 8.2 | 74 | 4600 | 100 | 8.2 | 28.9 | 8.2 |
| Selenium | 7782-49-2 | 71 | 290 | 4200 | 200 | 71 | 5 | 71 |
| Silver | 7440-22-4 | | 1.9 | + | | 0.23 | 0.12 | 1.9 |
| Thallium | 7440-28-0 | == | | 0.47 | 6.3 | 21.3 | 10 | 6.3 |
| Zinc | 7440-66-6 | 81 | 90 | 26000 | 1000 | 81 | 65.7 | 81 |
| Metals, Organic (ug/L) | • | | • | | | | | |
| Tributyltin | 688-73-3 | 0.0074 | 0.42 | ÷ | 22 | 0.001 (a) | i a | 0.0074 |
| Polycyclic Aromatic Hydrocarbons (PAHs) (ug/L) | • | | | | | | | |
| 1-Methylnaphthalene | 90-12-0 | : | | | . | 2.1 | | 2.1 |
| 2-Methylnaphthalene | 91-57-6 | | - | A-a-a- Sente | | 4.2 | 330 | 4.2 |
| Acenaphthene | 83-32-9 | | | 990 | 30 | 6.6 | 38 | 30 |
| Acenaphthylene | 208-96-8 | . n. | # |) | | | 4840 | 4840 |
| Anthracene | 120-12-7 | | - | 40000 | 100 | 0.18 | 0.035 | 100 |
| Benzo(a)anthracene | 56-55-3 | | | 0.018 | 0.00016 | 0.018 | 0.025 | 0.00016 |
| Benzo(a)pyrene | 50-32-8 | | = | 0.018 | 0.000016 | 0.015 | 0.014 | 0.000016 |
| Benzo(b)fluoranthene | 205-99-2 | H L | = | 0.018 | 0.00016 | = | 9.07 | 0.00016 |
| Benzo(b,k)fluoranthene | | 250 | | . | 0.0016 | 53 | .55 | 0.0016 |
| Benzo(g,h,i)perylene | 191-24-2 | == | | | | - | 7.64 | 7.64 |
| Benzo(j)fluoranthene | 205-82-3 | | | | | - | | == |
| Benzo(k)fluoranthene | 207-08-9 | 70/0 | | 0.018 | 0.0016 | | rest. | 0.0016 |
| Chrysene | 218-01-9 | | - | 0.018 | 0.016 | | | 0.016 |

Table 3-7 - Development of Initial PRGs for Surface Water

| | | | AR | ARs | | Other Screening B | enchmark Sources | |
|---|------------|--|--|---|---|--|--|---|
| Analyte | CAS Number | National Recommended Water Quality Criteria - Aquatic Life Criteria - Saltwater CCC (chronic) ¹ EPA, 2013 | National Recommended Water Quality Criteria - Aquatic Life Criteria - Saltwater CMC (acute) ¹ EPA, 2013 | National Recommended Water Quality Criteria - Human Health for the Consumption of Organisms EPA, 2013 | EPA 2016 CWA- Effective Human Health Criteria Applicable to Washington (Organism Only) EPA/Ecology,2016 | EPA Region 3 BTAG Marine Water Screening Benchmarks EPA, 2006 | EPA Region 5 RCRA - Ecological Screening Levels - Water EPA, 2003 | Initial PRGs used for Data Screening |
| Dibenzo(a,h)anthracene | 53-70-3 | | | 0.018 | 0.000016 | | | 0.000016 |
| Fluoranthene | 206-44-0 | | | 140 | 6 | 1.6 | 1.9 | 6 |
| Fluorene | 86-73-7 | | , | 5300 | 10 | 2.5 | 19 | 10 |
| Indeno(1,2,3-c,d)pyrene | 193-39-5 | | | 0.018 | 0.00016 | | 4.31 | 0.00016 |
| Naphthalene | 91-20-3 | | | = | | 1.4 (a) | 13 | 1.4 |
| Phenanthrene | 85-01-8 | | | = | | 1.5 | 3.6 | 1.5 |
| Pyrene | 129-00-0 | | - | 4000 | 8 | 0.24 | 0.3 | 8 |
| Total Benzofluoranthenes (b,j,k) | - | | | - | | | | (A) (A) |
| Total HPAH | | 1-1-1 | | (FE | | 45 | 2 4 | |
| Total LPAH | | | | | | | | |
| Total PAH | | | - | | | - | | |
| Polychlorinated Biphenyls (PCBs) (ug/L) | | | | | | | | |
| Aroclor 1016 | 12674-11-2 | | | ; | | | | |
| Aroclor 1221 | 11104-28-2 | | == | i | | | = | |
| Aroclor 1232 | 11141-16-5 | | == | i | | | | |
| Aroclor 1242 | 53469-21-9 | | | ; | | | | |
| Aroclor 1248 | 12672-29-6 | | | i | | | | |
| Aroclor 1254 | 11097-69-1 | | | ; | | - | | |
| Aroclor 1260 | 11096-82-5 | | | | | | | - |
| Aroclor 1262 | 37324-23-5 | | | | | | | |
| Aroclor 1268 | 11100-14-4 | | | | | - | | |
| Total PCB | | | == | - | 0.000007 | H- | | 0.000007 |
| Semivolatile Organic Carbons (SVOCs) (ug/L) | | | | | | | | |
| 1,2,4,5-Tetrachlorobenzene | 95-94-3 | | | 1.10 | | 129 | 3 | 1.10 |
| 1,2,4-Trichlorobenzene | 120-82-1 | | | 70 | 0.037 | 5.4 (a) | 30 | 0.037 |
| 1,2-Dichlorobenzene | 95-50-1 | | | 1300 | 800 | 42 (a) | 14 | 800 |
| 1,3-Dichlorobenzene | 541-73-1 | | | 960 | 2 | 28.5 | 38 | 2 |
| 1,4-Dichlorobenzene | 106-46-7 | (-1) | Ħ | 190 | 200 | 19.9 | 9.4 | 200 |
| 2,2'-Oxybis (1-chloropropane) | 108-60-1 | | | 65000 | 900 | - | | 900 |
| 2,3,4,6-Tetrachlorophenol | 58-90-2 | | | i | | 1.2 | 1.2 | 1.2 |
| 2,4,5-Trichlorophenol | 95-95-4 | | | l et | 200 400000 | 12 | | 12 |
| 2,4,6-Trichlorophenol | 88-06-2 | | - | 2.4 | 0.28 | 61 | 4.9 | 0.28 |
| 2,4-Dichlorophenol | 120-83-2 | | | 290 | 10 | 11 | 11 | 10 |
| 2,4-Dimethylphenol | 105-67-9 | | | 850 | 97 | | 100 | 97 |
| 2,4-Dinitrophenol | 51-28-5 | | == | 5300 | 100 | 48.5 | 19 | 100 |
| 2,4-Dinitrotoluene | 121-14-2 | 1-4 | = | 3.4 | 0.18 | 44 | 44 | 0.18 |
| 2,6-Dinitrotoluene | 606-20-2 | | | 1600 | 100 | 81 | 81 | 81 |
| 2-Chloronaphthalene | 91-58-7 | -55 | | 1600 | 100 | = | 0.396 | 100 |

Table 3-7 - Development of Initial PRGs for Surface Water

| | | | | AR | ARs | | Other Screening B | enchmark Sources | | |
|----------|---|------------|--|--|--|---|--|--|---|--|
| Analyte | | CAS Number | National Recommended Water Quality Criteria - Aquatic Life Criteria - Saltwater CCC (chronic) ¹ EPA, 2013 | National Recommended Water Quality Criteria - Aquatic Life Criteria - Saltwater CMC (acute) ¹ EPA, 2013 | National Recommended Water Quality Criteria - Human Health for the Consumption of Organisms EPA, 2013 | EPA 2016 CWA- Effective Human Health Criteria Applicable to Washington (Organism Only) EPA/Ecology,2016 | EPA Region 3 BTAG Marine Water Screening Benchmarks EPA, 2006 | EPA Region 5 RCRA - Ecological Screening Levels - Water EPA, 2003 | Initial PRGs used for Data Screening | |
| Allalyte | 2-Chlorophenol | 95-57-8 | | | 150 | 17 | 265 | 24 | 17 | |
| | 2-Methylphenol (o-Cresol) | 95-48-7 | | | | | 1020 | 67 | 67 | |
| | 2-Nitroaniline | 88-74-4 | 1-2 | | | 22 | | | | |
| | 2-Nitrophenol | 88-75-5 | | | | | 2940 | | 2940 | |
| | 3,3'-Dichlorobenzidine | 91-94-1 | | | 0.03 | 0.0033 | 73 | 4.5 | 0.0033 | |
| | 3-Methylphenol & 4-Methylphenol (m&p-Cresol) | 1319-77-3 | | = | | | | | | |
| | 3-Methylphenol (m-Cresol) | 108-39-4 | | | | | ===: | 62 | 62 | |
| | 3-Nitroaniline | 99-09-2 | | | | | - | | | |
| | 4-Bromophenyl-phenyl ether | 101-55-3 | | | | | 1.5 | 1.5 | 1.5 | |
| | 4-Chloro-3-methylphenol | 59-50-7 | | | | 36 | | 34.8 | 36 | |
| | 4-Chloroaniline | 106-47-8 | | | | | 232 | 232 | 232 | |
| | 4-Methylphenol (p-Cresol) | 106-47-8 | | | | | 543 | 25 | 25 | |
| | | 100-44-5 | | 10000 | Name of the last o | | | 1.000 | | |
| | 4-Nitroaniline | 100-01-6 | | | i | | | | | |
| | 4-Nitrophenol | 98-86-2 | | = | 14- | 44 | 71.7 | 60 | 60 | |
| | Acetophenone | 62-53-3 | | - | 5 | | 2.2 | | 2.2 | |
| | Aniline | | 14 | | | 24 | | 4.1 | | |
| | Atrazine | 1912-24-9 | | | <u></u> | | 1.8 | | 1.8 | |
| | Benzaldehyde | 100-52-7 | | | | | | | | |
| | Benzidine | 92-87-5 | H- | | 1 | 0.000023 | 3.9 | | 0.000023 | |
| | Benzoic acid | 65-85-0 | | | | | 42 | | 42 | |
| | Benzyl alcohol | 100-51-6 | | | | | 8.6 | 8.6 | 8.6 | |
| | Biphenyl (1,1'-Biphenyl) | 92-52-4 | | | | | 14 | | 14 | |
| | bis(2-Chloroethoxy)methane | 111-91-1 | 1-8 | | | | | | | |
| | bis(2-Chloroethyl)ether | 111-44-4 |) -1 | Ħ | 0.53 | 0.06 | == | 19000 | 0.06 | |
| | bis(2-Ethylhexyl)phthalate | 117-81-7 | | | 2.2 | 0.046 | 16 | 0.3 | 0.046 | |
| | Butylbenzyl phthalate | 85-68-7 | 1-2 | | 1900 | 0.013 | 29.4 | 23 | 0.013 | |
| | Caprolactam | 105-60-2 | 1-9 | | / | HH. | - | ;= | | |
| | Dibenzofuran | 132-64-9 | 1-1 | | | | 65 | 4 | 4 | |
| | Diethyl phthalate | 84-66-2 | | | 44000 | 200 | 75.9 | 110 | 200 | |
| | Dimethyl phthalate | 131-11-3 | | | 1100000 | 600 | 580 | | 600 | |
| | Di-n-butyl phthalate | 84-74-2 | | | 4500 | 8 | 3.4 | 9.7 | 8 | |
| | Dinitro-o-cresol (4,6-Dinitro-2-methylphenol) | 534-52-1 | 1-e | = | 280 | 7 | = | 23 | 7 | |
| | Di-n-octyl phthalate | 117-84-0 | | | | | 22 | 30 | 22 | |
| | Hexachlorobenzene | 118-74-1 | | - | 0.00029 | 0.00005 | 0.0003 | 0.0003 | 0.000005 | |
| | Hexachlorocyclopentadiene | 77-47-4 | | - | 1100 | 1 | 0.07 | 77 | 1 | |
| | Hexachloroethane | 67-72-1 | | | 3.3 | 0.02 | 9.4 | 8 | 0.02 | |
| | Isophorone | 78-59-1 | 1=9 | == | 960 | 110 | 129 | 920 | 110 | |
| | Nitrobenzene | 98-95-3 | 1-12 | | 690 | 100 | 66.8 | 220 | 100 | |

Table 3-7 - Development of Initial PRGs for Surface Water

| | | | AR | ARs | | Other Screening B | | |
|--|------------|-------------------------------|-------------------------------|-------------------------------|----------------------------------|----------------------|----------------------|----------------------------|
| | | National Recommended Water | National Recommended Water | National Recommended Water | EPA 2016 CWA- Effective Human | | | |
| | | Quality Criteria - | Quality Criteria - | Quality Criteria - | Health Criteria | | | |
| | | Aquatic Life Criteria - | Aquatic Life Criteria - | Human Health for the | Applicable to | EPA Region 3 BTAG | EPA Region 5 RCRA - | |
| | | Saltwater CCC | Saltwater CMC | Consumption of | Washington | Marine Water | Ecological Screening | |
| | | (chronic) ¹ | (acute) ¹ | Organisms | (Organism Only) | Screening Benchmarks | Levels - Water | Initial PRGs used for Data |
| Analyte | CAS Number | EPA, 2013 | EPA, 2013 | EPA, 2013 | EPA/Ecology,2016 | EPA, 2006 | EPA, 2003 | Screening |
| n-Nitrosodimethylamine | 62-75-9 | | | | 0.34 | 330000 | | 0.34 |
| n-Nitrosodi-n-propylamine | 621-64-7 | | | 0.51 | 0.058 | 120 | | 0.058 |
| n-Nitrosodiphenylamine | 86-30-6 | | | 6 | 0.69 | 33000 | H | 0.69 |
| Pentachlorophenol | 87-86-5 | 7.9 | 13 | 3 | 0.002 | 7.9 | 4 | 0.002 |
| Phenol | 108-95-2 | 1 55 | == | 860000 | 70000 | 58 | 180 | 70000 |
| Volatile Organic Carbons (VOCs) (ug/L) | | _ | | | | | | |
| 1,1,1-Trichloroethane | 71-55-6 | | = | | 50000 | 312 | 76 | 50000 |
| 1,1,1,2-Tetrachloroethane | 630-20-6 | 5 8 | | | | | | == |
| 1,1,2,2-Tetrachloroethane | 79-34-5 | | - | 4 | 0.3 | 90.2 | 380 | 0.3 |
| 1,1,2-Trichloroethane | 79-00-5 | | - | 16 | 0.9 | 550 | 500 | 0.9 |
| 1,1,2-Trichlorotrifluoroethane (Freon 113) | 76-13-1 | 14 | | | | | | |
| 1,1-Dichloroethane | 75-34-3 | | - | | - | 47 | 47 | 47 |
| 1,1-Dichloroethene | 75-35-4 | | - | 7100 | 4000 | 2240 | 65 | 4000 |
| 1,2,3-Trichlorobenzene | 87-61-6 | 1-0 | | + | Ŧ | 8 | H | 8 |
| 1,2,3-Trichloropropane | 96-18-4 | | - | | - | | | |
| 1,2,4-Trimethylbenzene | 95-63-6 | | # | | - | 19 | | 19 |
| 1,2-Dibromo-3-chloropropane | 96-12-8 | | - | | 1 | | | == |
| 1,2-Dichloroethane | 107-06-2 | | - | 37 | 73 | 1130 | 910 | 73 |
| 1,2-Dichloroethene, cis- | 156-59-2 | - | | - | I | | - | |
| 1,2-Dichloroethene, trans- | 156-60-5 | | | 10000 | 1000 | 970 | 970 | 1000 |
| 1,2-Dichloropropane | 78-87-5 | | - | 15 | 3.1 | 2400 | 360 | 3.1 |
| 1,3,5-Trimethylbenzene (Mesitylene) | 108-67-8 | | | | 1 | 71 | | 71 |
| 1,3-Dichloropropane | 142-28-9 | | | | I | | | - |
| 1,3-Dichloropropene, cis- | 10061-01-5 | 1-4 | | 21 | 1.2 | | H | 1.2 (b) |
| 1,3-Dichloropropene, trans- | 10061-02-6 | | - | 21 | 1.2 | | | 1.2 (b) |
| 1,4-Dichloro-2-butene, trans- | 110-57-6 | | - | | | | | |
| 1,4-Dioxane | 123-91-1 | 1-0 | | H- | Ŧ | | 22000 | 22000 |
| 2-Butanone (MEK) | 78-93-3 | | - | <u></u> | - | 14000 | 2200 | 2200 |
| 2-Chlorotoluene | 95-49-8 | 55 | - | | 1 | - | | |
| 2-Hexanone (Methyl butyl ketone) | 591-78-6 | | - | | 1 | 99 | 99 | 99 |
| 4-Chlorotoluene | 106-43-4 | | | | - | | | |
| 4-Isopropyltoluene (4-Cymene) | 99-87-6 | 144 | | | - | 85 | H | 85 |
| Acetone | 67-64-1 | | | 14- | | 564000 | 1700 | 1700 |
| Acrolein | 107-02-8 | | - | =- | 1.1 | 0.55 | 0.19 | 1.1 |
| Acrylonitrile | 107-13-1 | | - | | 0.028 | 581 | 66 | 0.028 |
| Benzene | 71-43-2 | | | 51.00 | 1.6 | 110 (a) | 114 | 1.6 |
| Bromobenzene | 108-86-1 | 144 | = | <u> </u> | - | = | 14 | |
| Bromochloromethane | 74-97-5 | | | | | | | |

Table 3-7 - Development of Initial PRGs for Surface Water

| | | | AR | ARs | | Other Screening B | enchmark Sources | |
|---|------------|---|---|--|---|---|---|--|
| | | National Recommended Water Quality Criteria - Aquatic Life Criteria - Saltwater CCC (chronic) ¹ | National Recommended Water Quality Criteria - Aquatic Life Criteria - Saltwater CMC (acute) ¹ | National Recommended Water Quality Criteria - Human Health for the Consumption of Organisms | EPA 2016 CWA- Effective Human Health Criteria Applicable to Washington (Organism Only) | EPA Region 3 BTAG Marine Water Screening Benchmarks | EPA Region 5 RCRA - Ecological Screening Levels - Water | Initial PRGs used for Data |
| Analyte | CAS Number | EPA, 2013 | EPA, 2013 | EPA, 2013 | EPA/Ecology,2016 | EPA, 2006 | EPA, 2003 | Screening |
| Bromodichloromethane | 75-27-4 | | | 17 | 2.8 | | | 2.8 |
| Bromoform (Tribromomethane) | 75-25-2 | | - | 140 | 12 | 640 | 230 | 12 |
| Bromomethane (Methyl bromide) | 74-83-9 | | | 1500 | 2400 | 120 | 16 | 2400 |
| Carbon disulfide | 75-15-0 | | - | <u></u> | | 0.92 | 15 | 0.92 |
| Carbon tetrachloride (Tetrachloromethane) | 56-23-5 | | | 1.6 | 0.35 | 1500 | 240 | 0.35 |
| Chlorobenzene | 108-90-7 | 144 | #4 | 1600 | 200 | 25 (a) | 47 | 200 |
| Chloroethane | 75-00-3 | 1 | | 1 | =- | | 1 | |
| Chloroform | 67-66-3 | 55 | | 470 | 600 | 815 | 140 | 600 |
| Chloromethane | 74-87-3 | | | | | 2700 | - | 2700 |
| Cyclohexane | 110-82-7 | | | | | 1 | | Name of the last o |
| Dibromochloromethane | 124-48-1 | 14 | ₩. | 13 | 2.2 | ## | H4 | 2.2 |
| Dibromomethane | 74-95-3 | | | 11- | | 1 | | |
| Dichlorodifluoromethane | 75-71-8 | | _ | | | _ | - | == |
| Dichloromethane (Methylene chloride) | 75-09-2 | 1 | - | 590 | 100 | 2560 | 940 | 100 |
| Ethylbenzene | 100-41-4 | | | 2100 | 31 | 25 (a) | 14 | 31 |
| Ethylene dibromide (1,2-Dibromoethane) | 106-93-4 | | | | | - | | == |
| Hexachlorobutadiene (Hexachloro-1,3-butadiene) | 87-68-3 | | | 18 | 0.01 | 0.3 | 0.053 | 0.01 |
| Isopropylbenzene (Cumene) | 98-82-8 | | | | | 2.6 | | 2.6 |
| Methyl acetate | 79-20-9 | 144 | н. | | ** | | | |
| Methyl iodide (Iodomethane) | 74-88-4 | | | ₩- | == | - | | |
| Methyl isobutyl ketone (4-Methyl-2-pentanone or (MIBK)) | 108-10-1 | == | | | | 123000 | 170 | 170 |
| Methyl tert-butyl ether (MTBE) | 1634-04-4 | | | | | 11070 | | 11070 |
| n-Butylbenzene | 104-51-8 | | | | | Andrew-weight News | | A TOPY OF THE PARTY OF THE PART |
| n-Propylbenzene | 103-65-1 | | | 14- | 22 | 128 | | 128 |
| o-Xylene | 95-47-6 | | | 1 | | | | |
| sec-Butylbenzene | 135-98-8 | | | | | | | |
| Styrene | 100-42-5 | F== | = | | == | 910 | 32 | 32 |
| tert-Butylbenzene | 98-06-6 | | | | | | | |
| Tetrachloroethene (PCE) | 127-18-4 | | | 3.3 | 2.9 | 45 | 45 | 2.9 |
| Toluene | 108-88-3 | | | 15000 | 130 | 215 (a) | 253 | 130 |
| Total xylene (reported, not calculated) | 1330-20-7 | | | | | 213 (d) | | |
| Total Xylene | | | | | | 19 | 27 | 19 |
| Trichloroethene (TCE) | 79-01-6 | 1 | | 30.00 | 0.7 | 21 | 47 | 0.7 |
| Trichlorofluoromethane (Fluorotrichloromethane) | 75-69-4 | | | | | | | |
| Vinyl acetate | 108-05-4 | | | | == | 16 | 248 | 16 |
| Vinyl acetate Vinyl chloride | 75-01-4 | | | 2.40 | 0.18 | 930 | 930 | 0.18 |

Table 3-7 - Development of Initial PRGs for Surface Water

Bremerton Gas Works Superfund Site

Bremerton, Washington

| | | | AR | ARs | | Other Screening B | | |
|---------|------------|-------------------------|-------------------------|----------------------|------------------------|----------------------|-----------------------------|----------------------------|
| | | | | | | | | |
| | | National | National | National | EPA 2016 CWA- | | | |
| | | Recommended Water | Recommended Water | Recommended Water | Effective Human | | | |
| | | Quality Criteria - | Quality Criteria - | Quality Criteria - | Health Criteria | | | |
| | | Aquatic Life Criteria - | Aquatic Life Criteria - | Human Health for the | Applicable to | EPA Region 3 BTAG | EPA Region 5 RCRA - | |
| | | Saltwater CCC | Saltwater CMC | Consumption of | Washington | Marine Water | Ecological Screening | |
| | | (chronic) ¹ | (acute) ¹ | Organisms | (Organism Only) | Screening Benchmarks | Levels - Water | Initial PRGs used for Data |
| Analyte | CAS Number | EPA, 2013 | EPA, 2013 | EPA, 2013 | EPA/Ecology,2016 | EPA, 2006 | EPA, 2003 | Screening |

Notes:

Compounds frequently associated with MGP-operations

'-- indicates not available

1 = Criteria for metals and methyl mercury are expressed in terms of the dissolved metal in the water column.

(a) = This is a Canadian Water Quality Guideline value and refers to the total concentration in an unfiltered sample.

(b) = 1,3-dichloropropene listed in EPA 2016 but it is not designated to the cis- or trans- isomers. Conservatively, the 1,3-dichloropropene screening value has been applied for initial evaluation.

BTAG = Biological Technical Assistance Group

CAS = Chemical Abstract Services

CCC = Criterion Continuous Concentration

CMC = Criterion Maximum Concentration

CWA = Clean Water Act

EPA = U.S. Environmental Protection Agency

HPAH = high molecular weight PAH

LPAH = low molecular weight PAH

L = liter

mg = milligram

MGP = manufactured gas plant

ng = nanogram

PRG = Preliminary Remediation Goal

RCRA = Resource Conservation and Recovery Act

RSL = regional screening level

μg = microgram

References:

EPA, 2003. EPA Region 5 Resource Conservation Recovery Act (RCRA) Ecological Screening Levels. August 22, 2003.

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EPA, 2013a. National Recommended Water Quality Criteria. Updated August 22, 2013. Available from:

http://water.epa.gov/scitech/swquidance/standards/criteria/current/index.cfm#altable.

EPA, 2016. EPAs Partial Approval/Partial Disapproval of Washington's Human Health Water Quality Criteria and Implementation

Tools. November 15, 2016. Available from: https://www.epa.gov/sites/production/files/2016-

11/documents/epas_partial_approvalpartial_disapproval_wa_hh_wqc_impl_tools_bellon_ltr_enclosures_508c.pdf

Table 3-8 - Summary of Data Quality Review for Existing Site Data

Bremerton Gas Works Superfund Site Bremerton, Washington

| | | | | Study/Media | | | |
|--|---|---|--|---|--|--|--|
| | 2008 E&E Targeted Brownfields (E&E 2008, E&E 2009 | 2010 E&E Removal Action (EPA 2010, AnchorQEA 2011) | 1995 Ecology (Ecology 1995) | 2007 Geoengineers (Geoengineers 2007a, 2007b) | 2008 E&E Targeted Brownfields (E&E 2008, E&E 2009) | 2007 Geoengineers (Geoengineers 2007a, 2007b) | 2008 E&E Targeted Brownfields (E&E 2008, E&E 2009) |
| Made Plan Description | Sediment | Sediment | Soil | Soil | Soil | Groundwater | Groundwater |
| Work Plan (SAP/QAPP) | Detailed QAPP covering multiple pieces of sampling program (soil, groundwater and sediment). Also includes general sediment sampling SOP and data report. | Site-Specific Sampling Plan (SSSP; not reviewed) approved by EPA, finalized after sampling conducted but in field deviations approved by EPA. | None | Work Plan, including site-specific SAP and QAPP, dated June 1, 2007 | SQAPP dated March 5, 2008 | Work Plan, including site-specific SAP and QAPP, dated June 1, 2007 | SQAPP dated March 5, 2008 |
| Collection methods and purpose | Detailed in QAPP. Sampling under EPA Brownfields management, follows EPA procedures. Limited for sediment; to determine if GW migration from upland sources is occurring into the Narrows. | Developed under EPA Superfund Technical Assessment Response Team (START). Determining origin of contamination from 12" exposed drain pipe on Sesko property beach. | Surface soil/sediment samples of suspected contamination based on visual inspection | Purpose to assess soil quality in potential contaminant source areas. Table of rationale for specific boring/sample locations referenced but not included in final work plan. | Judgmental sampling design to determine presence of contamination in areas of concern. Detailed rationale provided in SQAPP. | Purpose to assess groundwater quality in and downgradient of potential contaminant source areas. Table of rationale for specific boring/sample locations referenced but not included in final work plan. | Judgmental sampling design to determine presence of contamination in areas of concern. Detailed rationale provided in SQAPP. |
| Sample Location and Collection Methods | • | | | | 1 | | |
| Location method, accuracy, and datum. | Location established with GPS coordinates; accuracy not specified. Actual sampling appear to be close/at QAPP locations. Datum not specified. | Location established with GPS coordinates; accuracy not specified. Datum not specified. | Sample locations recorded on rough site sketch. No survey information provided. | Locations provided on scaled site map. Location method unknown. No survey information provided. | Locations provided on scaled site map. Location method unknown. No survey information provided. Note: locations of borings SP01 and SP03 apparently switched on site map, based on boring log information and correlation of chemical data with boring log observations. | | Locations provided on scaled site map. Location method unknown. No survey information provided. Note: locations of borings SP01 and SP03 apparently switched on site map, based on boring log information and correlation of chemical data with boring log observations. |
| Sample depths | 0-30cm | 0-6 inches | Less than 10 inches | up to 45 feet deep | up to 40 feet deep | 15-foot long well screens up to 45 feet deep | Monitoring Wells: 10-foot long well screens up to 45 feet deep. Temporary borings: depth not provided. |
| Collection method and matrix | Surface sediment. Dedicated stainless steel spoon. Collected at low tide from 5 biased locations targeted to evaluate potential for GW migration based on previous analytical and "on-site observations". | Surface sediment. Dedicated stainless steel spoon. Known areas of sediment deposition within the direct vicinity of the 12: drainpipe, collected below average high tide line. | Hand collection of surface soil/sediment samples | Hollow-stem auger drilling with split-spoon sampling. | Hollow-stem auger drilling with split-spoon sampling. | Report states low-flow sampling with peristaltic pump. Questionable for 30-ft deep groundwater samples. | Monitoring wells sampled using low-flow sampling using electric submersible pump. Methods for sampling temporary boreholes not provided. |
| Sample collection, processing and handling | Homogenized in dedicated stainless steel bowls (VOC cores taken from sampling locations prior to other sediment collection). Data report includes photographs at each sediment station. | Homogenized in dedicated stainless steel bowls (VOC cores taken from sampling locations prior to other sediment collection). Data report includes photographs at each sediment station. | Collection and handling activities not reported. | and field screened for contamination. 17 samples | Soil samples collected from 7 borings at 5-foot intervals and field screened for contamination. 48 samples collected for sample analysis. VOC samples collected by EPA 5035A. Protocols detailed in SAP. | Groundwater samples collected from 8 permanent, developed monitoring wells. Processing and handling protocols detailed in SAP. | Groundwater samples collected from 2 permanent, developed monitoring wells and 4 temporary borings. Processing and handling protocols detailed in SAP. |
| Holding time, preservation, and chain of custody | Detailed in the QAPP. Chain of custody provided in data report. Holding time and preservation discussed in lab data report. | Chain of custody provided in data report. Holding time and preservation discussed in lab data report. | chain of custody not provided. Laboratory case narrative indicates holding times were within recommended limits. | Requirements detailed in SAP and QAPP. Holding times and preservation were met as documented in data report. Chain of custody provided in data report. | Requirements detailed in SAP and QAPP. Holding times and preservation were met as documented in data report. Chain of custody provided in data report. | = | Requirements detailed in SAP and QAPP. Holding times and preservation were met as documented in data report. Chain of custody provided in data report. |
| Laboratory Analysis | | | | | | | |
| Analytical methods are standard or USEPA approved | EPA and NWTPH methods. TPH-Dx, TPH-Dx, VOC, SVOC, TAL metals. | EPA methods. VOC by 8260, SVOC by 8270, static sheen test. | EPA Methods. Metals - EPA200.7, EPA270.2, EPA206.2, EPA279.2, EPA245.5 PAHs - Manchester Modification of SW8270 | EPA and NWTPH Methods. TPH - Ecology NWTPH-Gx and NWTPH-Dx VOCs - EPA -8260B SVOCs - EPA 8270 SIM PCBs - EPA 8082 PP metals/chromiumVI - EPA 6000/7000 series TBT - Krone (GC/MS) | TPH - Ecology NWTPH-Gx and NWTPH-Dx VOCs - EPA 8260B SVOCs - EPA 8270C TAL metals - EPA 6000/7000 series | EPA and NWTPH Methods. TPH - Ecology NWTPH-Gx and NWTPH-Dx VOCs - EPA -8260B SVOCs - EPA 8270 SIM PCBs - EPA 8082 PP metals/chromiumVI - EPA 6000/7000 series | EPA and NWTPH Methods. TPH - Ecology NWTPH-Gx and NWTPH-Dx VOCs - EPA 8260B SVOCs - EPA 8270C TAL metals - EPA 6000/7000 series |
| Detection limits and qualifiers determined based on USEPA guidance | Yes. Detailed in the QAPP. Qualifier identified in laboratory data report. | Yes. Qualifier identified in laboratory data report. | summarized in QA narrative in laboratory data report. | Yes. Detailed in QAPP. Qualifiers identified in laboratory data report. | Yes. Detailed in QAPP. Qualifiers identified in laboratory data report. | Yes. Detailed in QAPP. Qualifiers identified in laboratory data report. | Yes. Detailed in QAPP. Qualifiers identified in laboratory data report. |
| Measurement instruments and calibration procedures | Detailed in QAPP. Sampling under EPA Brownfields management, follows EPA procedures. | Some detail provided in data validation memo. | Some detail provided in QA narrative in laboratory data report. | Yes. Detailed in QAPP. | Detailed in QAPP. Sampling under EPA Brownfields management, follows EPA procedures. | Yes. Detailed in QAPP. | Detailed in QAPP. Sampling under EPA Brownfields management, follows EPA procedures. |
| Quality Control and Data Validation | | | | | | | |
| Field/Lab quality control samples (duplicates, blanks) | Field rinsate and trip blanks (no issues in sediment samples) MS/MSD, serial dilution, internal standards. | Field trip blank. | MS/MSD, LCS | Field duplicate; method blanks, calibration blanks, sample blanks, MS/MSD, and LCS. | Laboratory blanks, rinsate blanks, trip blanks, MS/MSD. | Field duplicate, rinseate blank, and trip blanks; method blanks, calibration blanks, sample blanks, MS/MSD, and LCS. | Laboratory blanks, rinsate blanks, trip blanks, MS/MSD. |
| Analytical chemistry data must have been validated and qualified consistent with EPA functional guidelines | Data validation conducted. Data validation memo included as Appendix to data report. Procedures also detailed in QAPP. | Data validation conducted. Data validation memo included as Appendix to data report. | QA summary by lab. Compounds with low matrix spike recoveries rejected or "J" qualified. | QA summary by lab. | QA/QC review and data validation documented in data report. | QA summary by lab. | QA/QC review and data validation documented in data report. |
| Laboratory data reports | Level II Data Package Available. | Level II Data Package Available. | Partial Level II Data Package Available. | Level II Data Package Available | Level II Data Package Available | Level II Data Package Available | Level II Data Package Available |

COC = chemical of concern EPA = U.S. Environmental Protection Agency GC/MS = gas chromatography-mass spectrometry LCS = laboratory control sample MS/MSD = matrix spike/matrix spike duplicate NWTPH = Northwest total petroleum hydrocarbon PAH = polycyclic aromatic hydrocarbon PCB = polychlorinated biphenyl PP = priority pollutant QA = quality assurance QAPP = Quality Assurance Project Plan QC = quality control SAP = Sampling and Analysis Plan SOP = standard operating procedure SQAPP = SAP/QAPP SVOC = semivolatile organic compound TAL = target analyte list TBT = tributyltin TPH = total petroleum hydrocarbons

Washington State Department of Ecology (Ecology), 1995, Initial Investigation Inspection, Sesko Property, March 29, 1995.

GeoEngineers, 2007a, Preliminary Upland Assessment Work Plan, McConkey/Sesko Site, June 1, 2007.

GeoEngineers, 2007b, Preliminary Upland Assessment Report, McConkey/Sesko Brownfield Site, Prepared by GeoEngineers, Inc., for the City of Bremerton, October 26, 2007.

Ecology & Environment, Inc. (E&E), 2008, Final Bremerton Gasworks Targeted Brownfields Assessment Sampling and Quality Assurance Project Plan, Prepared by E&E for EPA, March 5, 2008.

Ecology & Environment, Inc. (E&E), 2009, Final Bremerton Gasworks Targeted Brownfields Assessment Report, Prepared by E&E for U.S. Environmental Protection Agency, August 2009.

Anchor QEA, 2011, Final Completion Report: Former Bremerton MGP Site, Incident Action and Time Critical Removal Action, Prepared for U.S. Coast Guard Sector Puget Sound Incident Management Division on behalf of Cascade Natural Gas Corporation, January 2011.

In Proceedings of the Cascade Natural Gas Corporation, January 2011.

US Environmental Protection Agency (EPA), 2010 Site-Specific Sampling Plan, Bremerton MGP Release, October 28, 2010.

VOC = volatile organic compound

Table 3-9 - Summary of Data Quality Review for Existing Sediment and Tissue Data

Bremerton Gas Works Superfund Site Bremerton, WA

| | 2010 and 2012 ENVVEST | 2005, 2007 NOAA Mussel Watch @ station | Study/Media | 2008-2009 PSAMP - Spatial/Temporal - Central | 1989-2013 PSAMP Long term/ temporal | 2009 - PSAMP Urban Waters Initiative - Bainbridge |
|--|---|--|--|---|---|--|
| | 2010 and 2012 ENVVES1 | SIWP | 2001 303d Ecology Clam Crab | Sound | 1989-2013 PSAMP Long term/ temporal | Basin |
| | Mussel tissue. Data from 11 locations in Dyes Inlet and Sinclair Inlet considered for regional information. | Mussel Tissue. Data from 1 location in Sinclair Inlet considered for regional information. | Clam and crab tissue. Data from 3 locations in Dyes Inlet considered for regional information. | Sediment. Data from 11 locations in Dyes Inlet and Sinclair Inlet considered for regional information. | Sediment. Data from 1 location in Sinclair Inlet considered for regional information. | Sediment. Data from 18 locations in Dyes Inlet and Sinclair Inlet considered for regional information. |
| Work Plan Documentation | | | | | | |
| Work Plan (SAP/QAPP) | Detailed SAP/QAPP developed with EPA and Ecology under the cooperative Environmental Investment (ENVVEST) program (Johnston et al. 2009; 2010). | Detailed SAP/QAPP developed under NOAA National Status and Trends Program (NOAA 1993 and 2006). | Ecology (2001) QAPP. Results summarized in the 2002 data report and queried from EIM. | Detailed programattic CAPP (2009) developed cooperatively with State and Federal agencies. Event-specific addenda (2010, 2011, 2012). | Detailed programattic QAPP (2009) developed cooperatively with State and Federal agencies. Event-specific addenda (2010, 2011, 2012). | Detailed programattic QAPP (2009) developed cooperatively with State and Federal agencies. Event-specific addenda (2010, 2011, 2012) . |
| Collection methods, purpose and representativeness | Hand collection of blue mussels (Mytilus spp.) via boat or from shore. Shucked, whole organism. Methods follow NOAA protocol. Location control details provided. | 1 | Hand collection of male cancer crab tissue (Cancer gracilis) via crab pots (though Dungeness and Blue crabs targeted but none found); native and Japanese little neck clam tissue via hand digging (Protothaca staminea and Tapes japonica). | | 0.1 m2 modified stainless steel van Veen, lowered via cable to open upon sediment contact. Targeted fine grained sediment, sample rejected in field if not fine-grained dominant during in-field visual inspection. | 0.1 m2 modified stainless steel van Veen, lowered via cable to open upon sediment contact. Targeted fine grained sediment, sample rejected in field if no fine-grained dominant during in-field visual inspection. |
| Sample Location and Collection Methods | | | | | | |
| Location method, accuracy and datum | Location established with GPS; accuracy not specified. Table provided with coordinates. Datum not specified. | Location established with GPS. Accuracy and datum not specified. | Location established with GPS, accuracy not specified. Table provided with coordinates. Datum is NAD 83. | Location established with differential GPS. with expected accuracy of better than 3 meters. Table provided with coordinates. Datum is NAD 83. | Location established with differential GPS. with expected accuracy of better than 3 meters. Table provided with coordinates. Datum is NAD 83. | Location established with differential GPS. with expected accuracy of better than 3 meters. Table provided with coordinates. Datum is NAD 83. |
| Sample depths | Above MLLW - on rocks, piling, cabling, piers. | Detailed in NOAA (1993) SAP. Depends on station, some shoreline, some underwater. | Crabs: via pots on surface Clams: via hand digging within 100 sq ft of beach. | Top 2-3cm. | Top 2-3cm. | Top 2-3cm. |
| Sample collection, processing and handling | Field - Hand harvest, cut byssus threads with knife; hand brush off debris; 1-3 replicates per stations (reps within 150' radius of station loc; 30-50 mussels per replicate. Hand delivery to lab. <u>Lab</u> - kept at -20C until measured and shucked with ceramic knife; rinsed with DI, composite by replicate then by station using Ti blender. | Field - Detailed in NOAA (1993) SAP. In general, some stations hand collection or with rake, some with bivalve dredge. Lab - shell size and volume determined; shucked; homogenized using stainless steel blender with titanium blades. Chemically dried using hydromatrix. | Detailed in SAP.Crabs: Muscle tissue (no organs or | Field - stainless steel spoon from each grab; grabs composited into stainless steel bucket; salinity and sediment temp measured. | Field - stainless steel spoon from each grab; grabs composited into stainless steel bucket; salinity and sediment temp measured. | Field - stainless steel spoon from each grab; grabs composited into stainless steel bucket; salinity and sediment temp measured. |
| Holding time, preservation, and chain of custody | Requirements detailed in the QAPP. Holding times and preservation were met as documented in the data report. Chain of custody provided in the data report. | | Requirements detailed in the QAPP. Holding times and preservation were met as documented in the data report. Chain of custody provided in the data report. | Requirements detailed in the QAPP. Holding times and preservation were met as documented in the data report. Chain of custody provided in the data report. | Requirements detailed in the QAPP. Holding times and preservation were met as documented in the data report. Chain of custody provided in the data report. | Requirements detailed in the QAPP. Holding times and preservation were met as documented in the data report. Chain of custody provided in the data report. |
| Laboratory Analysis | | | 1 | | 1 | 1 |
| Analytical methods are standard or EPA approved | Total Hg - EPA 7473m (EPA 1631 rev E in QAPP). Battelle SOPS for other metals and PCB congeners, PAHs - GC/MS Battelle SOP -015. Standard analytical methods. Lipids, moisture, C and N isotopes, trace metals, Hg, isotopes, 20 NS&T PCB congeners, parent and alkylated PAH. | Lipids, moisture, C and N isotopes, trace metals, Hg, isotopes, 20 NS&T PCB congeners, parent and alkylated PAH. Detailed in specific analytical methods reports. Standard analytical methods. | Lipid, and imony, SVOCs, PAHs. USEPA and PSEP standard anlytical methods. | Grain size, TOC, metals, pesticides, chlorobenzenes, PAHs, phenolics, phthalates, PCBs, PBDEs, bPA, triclosan, and other misc. including HCBD, dibenzofuran, carbazole and tin. EPA and PSEP standard analytical methods. | USEPA and PSEP standard analytical methods. | USEPA and PSEP standard analytical methods. |
| Detection limits and qualifiers determined based on EPA guidance | Yes. Detailed in QAPP and summarized in QA/QC narrative in data report. | Yes. Detailed in QAPP and summarized in QA/QC narrative in data report. | Yes. Detailed in QAPP and summarized in QA/QC narrative in data report. | Yes. Detailed in QAPP and summarized in QA/QC narrative in data report. | Yes. Detailed in QAPP and summarized in QA/QC narrative in data report. | Yes. Detailed in QAPP and summarized in QA/QC narrative in data report. |
| Measurement instruments and calibration procedures | Detailed in QAPP. | Detailed in QAPP. | Detailed in QAPP. | Detailed in QAPP. | Detailed in QAPP. | Detailed in QAPP. |
| Quality Control and Data Validation | | | l | | · | I |
| Field/Lab quality control samples (duplicates, blanks) | B, BS, MS/MSD, LD, reference material. | B, BS, MS/MSD, LD, reference material. | Blank, MS/MSD. | Blind field split replicates, field blanks; lab replicates, MS/MSD, lab control, MB, reference material. | Blind field split replicates, field blanks; lab replicates, MS/MSD, lab control, MB, reference material. | Blind field split replicates, field blanks; lab replicates, MS/MSD, lab control, MB, reference material. |
| Analytical chemistry data must have been validated and qualified consistent with EPA functional guidelines | Data validation conducted. Details in case narratives. | Not available online. | Data validation conducted. Details in case narratives. | Data validation conducted. Details in case narratives. | Data validation conducted. Details in case narratives. | Data validation conducted. Details in case narratives. |
| Laboratory data reports | Level II Data Package Available. | Not available online. | Case narrative text only. | Level II Data Package Available. | Only case narratives available through 2000. Online archives incomplete. | Level II Data Package Available. |

B = Blank bPA = Bisphenol A BS = Blank spike COCs = chemical of concerns EPA = U.S. Environmental Protection Agency HCBD = Hexachlorobutadiene GPS = global positioning system LCS = Laboratory control sample MB = Method blank MS/MSD = Matrix spike/matrix spike duplicate MLLW = Mean lower-low water NOAA = National Oceanic and Atmospheric Administration

PAHs = polycyclic aromatic hydrocarbons PBDE = Polybrominated diphenyl ether

PCBs = polychlorinated biphenyls PSEP = Puget Sound Estuary Program QAPP = Quality Assurance Project Plan QA/QC = quality assurance/quality control

SAP = Sampling and Analysis Plan SVOC = semivolatile organic compound TAL = Target analyte list TOC = Total organic carbon TPH = total petroleum hydrocarbons VOC = volatile organic compound

1989-2013 PSAMP 2008-2009 PSAMP 2009 PSAMP

2010 and 2012 ENVVEST (Johnston 2010 and Brandenberger 2012)

Johnston et al. 2009; 2010

2005, 2007 NOAA Mussel Watch

2001 303d Ecology Clam Crab

PSAMP. 2009. Quality Assurance Project Plan. The Puget Sound Assessment and Monitoring Program: Sediment Monitoring Component. August 2009. Publication No. 09-03-12: PSAMP. 2010 Addendum to Quality Assurance Project Plan. The Puget Sound Assessment and Monitoring Program: Sediment Monitoring Component. August 2010. Publication No. 09-03-121-Addendum: PSAMP. 2011 Addendum to Quality Assurance Project Plan. The Puget Sound Assessment and Monitoring Program: Sediment Monitoring Component. August 2010. Publication No. 09-03-121-Addendum

PSAMP. 2012 Addendum to Quality Assurance Project Plan. The Puget Sound Assessment and Monitoring Program/Urban Waters Initiative: Sediment Monitoring in the San Juan Islands and Port Gardner/ Everett Harbor. December 2011. Publication No. 09-03-121-Addendum Brandenberger JM, CR Suslick, LI Kuo RK Johnston. 2012. Ambient Monitoring for Sinclair and Dyes Inlets, Puget Sound, Washington: Chemical Analyses for 2012 Regional Mussel Watch. Prepared for the U.S. Department of Energy. PNNL-21862. September 201

Striplin, P.L., 1988. Puget Sound Ambient Monitoring Program: Marine Sediment Quality Implementation Plan. Washington State Department of Ecology, Olympia, Washington. 57 pp. www.ecy.wa.gov/biblio/88e37.html. Also see QAPP addendum PSAMP (2009, 2010, 2011, and 2012)

Johnston, RK, GH Rosen, JM Bandenberger, J.M. Wright, E. Mollerstuen, J. Young, and T. Tompkins. 2010. Sampling and Analysis Plan for Ambient Monitoring and Toxicity Testing for Sinclair and Dyes Inlets, Puget Sound, Washington. Prepared for Project ENVVEST, Puget Sound Naval Shipyard & Intermediate Maintenance

Facility, Bremerton, WA, Revised Sept. 18, 2010. Johnston, R.K., G.H. Rosen, J.M. Brandenberger, V.S. Whitney, and J.M. Wright. 2009. Sampling and Analysis Plan for Ambient Monitoring and Toxicity Testing for Sinclair and Dyes Inlets, Puget Sound, Washington. ENVVEST Planning Document National Oceanic and Atmospheric Adiministration (NOAA). 1993. Sampling and Analytical Methods of the National Status and Trends Program National Benthic Surveillance and Mussel Watch Projects 1984-1992. Volumes I through IV. NOAA Technical Memorandum NOS ORCA 71. G. G. Lauenstein and A. Y. Cantillo (Editors

NOAA. 2006a. Kimbrough, K. L., and G. G. Lauenstein (Editors). 2006. Major and Trace Element Analytical Methods of the National Status and Trends Program: 2000-2006. Silver Spring, MD. NOAA Technical Memorandum NOS NCCOS 29, 19 pt NOAA. 2006b. Kimbrough, K. L., G. G. Lauenstein and W. E. Johnston (Editors). 2006. Organic Contaminant Analytical Methods of the National Status and Trends Program: Update 2000-2006. NOAA Technical Memorandum NOS NCCOS 30, 137 pt

NOAA. 2008. Kimbrough, K. L., W. E. Johnston, G. G. Lauenstein, J. D. Christensen and D. A. Apeti.. An Assessment of Two Decades of Contaminant Monitoring in the Nation's Coastal Zone. Silver Spring, MD. NOAA Technical. Memorandum NOS NCCOS 74. 105 pp.

Washington State Department of Ecology (Ecology). 2002. Results of Sampling to Verify 303(d) Listings for Chemical Contaminants in Shellfish from Dyes Inlet and Port Washington Narrows. March 2002. Publication No. 02-03-01

Table 3-10 - Summary of Soil DataBremerton Gas Works Superfund Site
Bremerton, Washington

| Total Petroleum Hydrocarbons (TPH) | Gasoline Range Hydrocarbons Diesel Range Hydrocarbons Oil Range Hydrocarbons Aluminum Antimony Arsenic Barium Beryllium Cadmium Calcium Chromium (Total) Chromium (VI) Cobalt Copper Iron Lead Magnesium | 15 15 15 7 13 15 7 15 7 15 15 7 | 59 58 58 42 31 59 42 59 59 | 11 10 11 42 2 59 42 | 645 36,000 29,000 24,100 1.2 48.4 | 5 17.1 18 5,780 | | | | | | |
|---|--|--|--|---------------------------------------|--|--------------------------|--|----------|----|--------|---|--|
| Total Petroleum Hydrocarbons (TPH) | Diesel Range Hydrocarbons Oil Range Hydrocarbons Aluminum Antimony Arsenic Barium Beryllium Cadmium Calcium Chromium (Total) Chromium (VI) Cobalt Copper Iron Lead | 15 7 13 15 7 15 15 15 7 | 58 42 31 59 42 59 59 42 | 11 42 2 59 42 | 36,000 29,000 24,100 1.2 | 18 5,780 | | | | | | |
| | Oil Range Hydrocarbons Aluminum Antimony Arsenic Barium Beryllium Cadmium Calcium Chromium (Total) Chromium (VI) Cobalt Copper Iron Lead | 7 13 15 7 15 15 15 7 | 42 31 59 42 59 59 59 | 42 2 59 42 | 29,000 24,100 1.2 | 18 5,780 | | | | | i e e e e e e e e e e e e e e e e e e e | |
| | Aluminum Antimony Arsenic Barium Beryllium Cadmium Calcium Chromium (Total) Chromium (VI) Cobalt Copper Iron Lead | 15 7 15 15 7 15 | 31 59 42 59 59 42 | 2 59 42 | 1.2 | | | i l | | | | |
| Metals | Arsenic Barium Beryllium Cadmium Calcium Chromium (Total) Chromium (VI) Cobalt Copper Iron Lead | 15 7 15 15 7 15 | 59 42 59 59 42 | 59 42 | | | 7,700 | 34 | | 32,600 | | |
| Metals | Barium Beryllium Cadmium Calcium Chromium (Total) Chromium (VI) Cobalt Copper Iron Lead | 7 15 15 7 15 | 42 59 59 42 | 42 | 48.4 | 0.8 | 0.27 | 2 | 29 | 5 | | 12 |
| Metals | Beryllium Cadmium Calcium Chromium (Total) Chromium (VI) Cobalt Copper Iron Lead | 15 7 15 | 59 59 42 | | | 0.5 | 0.68 | 55 | | 7 | 2 | |
| Metals | Cadmium Calcium Chromium (Total) Chromium (VI) Cobalt Copper Iron Lead | 15 7 15 | 59 42 | | 120 | 23.9 | 330 | | | 255 | | |
| Metals | Calcium Chromium (Total) Chromium (VI) Cobalt Copper Iron Lead | 7 15 | 42 | 42 | 0.5 | 0.1 | 16 | 0.4 | 05 | 0.6 | | 7 |
| Metals | Chromium (Total) Chromium (VI) Cobalt Copper Iron Lead | | | 34 | 1.6 | 0.2 | 0.36 | 21 | 25 | 1 | 4 | |
| Metals | Chromium (VI) Cobalt Copper Iron Lead | | E0. | 42 59 | 21,300 60.8 | 1,620 | 26 | 22 | | 48 | 11 | |
| Metals | Cobalt Copper Iron Lead | 7 | 59 17 | 59 | 60.8 | 14.6 | 26 0.3 | 32 | 17 | 46 | 11 | |
| Metals ———————————————————————————————————— | Copper Iron Lead | | 42 | 42 | 19 | 3.3 | 2.3 | 42 | 17 | 11 | 15 | |
| Metals ———————————————————————————————————— | Iron Lead | 15 | 59 | 59 | 79.1 | 8 | 2.3 | 18 | | 36 | 17 | |
| Metals | Lead | 7 | 42 | 42 | 47,800 | 9,570 | 5,500 | 42 | | 36,100 | 3 | |
| | | 15 | 59 | 57 | 246 | 0.6 | 11 | 7 | | 24 | 6 | |
| | | 7 | 42 | 42 | 14,900 | 1,380 | - | <u>'</u> | | | <u> </u> | |
| | Manganese | 7 | 42 | 42 | 824 | 170 | 180 | 38 | | 1,200 | | |
| E | Mercury | 15 | 59 | 14 | 1.62 | 0.1 | 1.1 | 1 | | 0.07 | 14 | 45 |
| | Nickel | 15 | 59 | 59 | 66.3 | 21.2 | 38 | 27 | | 48 | 17 | |
| _ | Potassium | 7 | 42 | 42 | 2,000 | 233 | | | | | | |
| | Selenium | 15 | 59 | | · | | 0.52 | | 57 | 0.78 | | 42 |
| | Silver | 15 | 59 | | | | 4.2 | | | 0.61 | | 46 |
| | Sodium | 7 | 42 | 42 | 565 | 120 | | | | | | |
| | Thallium | 15 | 59 | 34 | 5.7 | 1.1 | 0.078 | 34 | 25 | | | |
| | Vanadium | 7 | 42 | 42 | 86 | 20.7 | 7.8 | 42 | | 45 | 17 | |
| | Zinc | 15 | 59 | 59 | 376 | 18.9 | 46 | 23 | | 85 | 5 | |
| | Acenaphthene | 18 | 60 | 19 | 31.2 | 0.0012 | 360 | | | | | |
| <u> </u> | Acenaphthylene | 23 20 | 61 | 23 | 460 274 | 0.00091 0.0012 | 1800 | | | | | |
| <u> </u> | Anthracene | 19 | 61 61 | 24 46 | 79 | 0.0012 | 1000 | | | | | |
| <u> </u> | Benzo(g,h,i)perylene Dibenzofuran | 15 | 59 | 46 | 0.37 | 0.00071 | 7.3 | | 3 | | | |
| | Fluoranthene | 22 | 61 | 32 | 572 | 0.00068 | 240 | 1 | 3 | | | |
| Polycyclic Aromatic Hydrocarbons (PAHs) | Fluorene | 20 | 61 | 25 | 404 | 0.0007 | 240 | 1 | | | | |
| | Phenanthrene | 24 | 61 | 39 | 1490 | 0.00061 | 1 2.10 | ' | | | | |
| | Pyrene | 21 | 61 | 38 | 913 | 0.0006 | 180 | 2 | | | | |
| | 1-Methylnaphthalene | 12 | 17 | 10 | 615 | 0.0144 | 18 | 5 | | | | |
| | 2-Methylnaphthalene | 13 | 17 | 10 | 978 | 0.0158 | 24 | 5 | | | | |
| | Naphthalene | 10 | 12 | 11 | 953 | 0.00047 | 3.8 | 4 | | | | |
| | Benz(a)anthracene | 18 | 61 | 29 | 113 | 0.0011 | 0.16 | 15 | 2 | | | |
| | Benzo(a)pyrene | 17 | 61 | 40 | 116 | 0.00053 | 0.016 | 21 | | | | |
| <u> </u> | Benzo(b)fluoranthene | 17 | 61 | 29 | 57.4 | 0.00085 | 0.16 | 15 | 1 | | | |
| <u> </u> | Benzo(k)fluoranthene | 17 | 61 | 36 35 | 60.6 | 0.00056 | 1.6 | 10 | | | | |
| <u> </u> | Chrysene Dibenzo(a,h)anthracene | 17 17 | 61 61 | 35 36 | 146 22.8 | 0.00067 0.0008 | 16 0.016 | 6 16 | 3 | | | |
| Carcinogenic PAHs (cPAHs) | Indeno(1,2,3-cd)pyrene | 17 | 61 | 44 | 58.5 | 0.0006 | 0.016 | 15 | 1 | | | |
| - | Total cPAHs TEQ (ND = 0) | 17 | 61 | 50 | 149 | 0.000066 | 0.16 | 21 | ' | | | |
| | otal cPAHs TEQ (ND = 1/2 RDL) | 17 | 61 | 50 | 149 | 0.000842 | 0.016 | 21 | | | | |
| 100 | Total HPAHs | 2 | 2 | 2 | 76.22 | 21.32 | 1.1 | 2 | | | | |
| ⊢ | Total LPAHs | 2 | 2 | 2 | 121.5 | 3.813 | 29 | 1 | | | | |
| | Total PAHs | 2 | 2 | 2 | 197.72 | 25.133 | | · | | | | |
| | 1,1'-Biphenyl | 7 | 42 | 5 | 0.98 | 0.014 | 4.7 | | | | | |
| | 1,2,4,5-Tetrachlorobenzene | 7 | 42 | | | | 2.3 | | | | | |
| | 1,2-Dichlorobenzene | 1 | 1 | | | | 180 | | | | | |
| | 1,3-Dichlorobenzene | 7 | 40 | | | | | | | | | |
| | 1,4-Dichlorobenzene | 1 | 1 | | | | 2.6 | | | | | |
| | 1,4-Dioxane | 7 | 42 | | | | 5.3 | | 2 | | | |
| | 2,3,4,6-Tetrachlorophenol | 7 | 42 | | | | 190 | | | | | |
| | 2,4,5-Trichlorophenol | 15 | 59 | | | | 630 | | | | | |
| <u> </u> | 2,4,6-Trichlorophenol | 15 | 59 | | | | 6.3 | | 4 | | | |
| <u> </u> | 2,4-Dichlorophenol 2,4-Dimethylphenol | 15 15 | 59 59 | | 0.031 | 0.031 | 19 | 1 | 2 | | - | i |

Table 3-10 - Summary of Soil DataBremerton Gas Works Superfund Site
Bremerton, Washington

| Chemical Group | Chemical Constituent | Number of Locations | Number of Samples | Number of Detections | Maximum Detected Concentration (mg/kg) | Minimum Detected Concentration (mg/kg) | Soil PRG (mg/kg) | Detected Concentrations Exceeding the PRG | Number of Non- Detect Results with Reporting Limit Concentrations that Exceed the PRG | Puget Sound Background Metals Concentration (mg/kg)1 | Number of Detected Concentrations Exceeding Puget Sound Background Concentration | Number of Non- Detect Results with Reporting Limit Concentrations that Exceed Puget Sound Background Concentration |
|--------------------------------------|--|------------------------|----------------------|-------------------------|---|---|---------------------|--|---|--|--|--|
| Gridina Group | 2,4-Dinitrophenol | 15 | 59 | Dottootiono | (9,9) | (9/9/ | 13 | | 3 | (9/1.9/ | Concontaction | |
| | 2-Chloronaphthalene | 15 | 59 | | | | 480 | | <u> </u> | | | |
| | 2-Chlorophenol | 15 | 59 | | | | 39 | | 2 | | | |
| | 2-Methylphenol | 8 | 17 | | | | 320 | | | | | |
| | 2-Nitroaniline | 15 | 59 | | | | 63 | | 2 | | | |
| | 2-Nitrophenol | 15 | 59 | | | | | | | | | |
| | 3 & 4 Methylphenol | 8 | 17 | | | | 4.0 | | 47 | | | |
| | 3,3'-Dichlorobenzidine 3-Nitroaniline | 15 15 | 59 59 | | | | 1.2 | | 17 | | | |
| | 4,6-Dinitro-2-methylphenol | 15 | 59 | | | | 0.51 | | 17 | | | |
| | 4-Bromophenyl phenyl ether | 15 | 59 | | | | 0.01 | | 17 | | | |
| | 4-Chloro-3-methylphenol | 15 | 59 | | | | 630 | | | | | |
| | 4-Chloroaniline | 15 | 59 | | | | 2.7 | | 8 | | | |
| | 4-Chlorophenyl phenyl ether | 15 | 59 | | | | | | | | | |
| | 4-Methylphenol | 7 | 42 | | | | 630 | | | | | |
| | 4-Nitroaniline | 15 | 59 50 | | | | 25 | | 2 | | | |
| | 4-Nitrophenol Acenaphthene | 15 | 59 1 | | | | 360 | 1 | | | | |
| | Acetophenone Acetophenone | 7 | 42 | 2 | 1.5 | 0.03 | 300 | | | | | |
| | Aniline | 8 | 17 | | 1.0 | 0.00 | 44 | | 2 | | | |
| Other Semi-Volaile Organic Compounds | Atrazine | 7 | 42 | | | | | | - | | | |
| (SVOCs) | Benzaldehyde | 7 | 42 | | | | | | | | | |
| | Benzidine | 7 | 42 | | | | 0.00053 | | 42 | | | |
| | Benzoic acid | 8 | 17 | | | | 25,000 | | | | | |
| | Benzyl alcohol | 8 | 17 | _ | 2 200 | 0.045 | 630 | | | | | |
| | Benzyl butyl phthalate Bis(2-chloro-1-methylethyl) ether | 15 15 | 59 59 | 5 | 0.029 | 0.015 | 290 310 | | | | | |
| | Bis(2-chloroethoxy)methane | 15 | 59 | | | | 19 | | 2 | | | |
| | Bis(2-chloroethyl) ether | 15 | 59 | | | | 0.23 | | 17 | | | |
| | Bis(2-ethylhexyl) phthalate | 15 | 59 | 39 | 0.29 | 0.069 | 39 | | 2 | | | |
| | Caprolactam | 7 | 42 | 1 | 0.015 | 0.015 | | | | | | |
| | Carbazole | 15 | 59 | 5 | 0.49 | 0.019 | | | | | | |
| | Dibenzofuran | 2 | 2 | 2 | 2.6 | 0.082 | 7.3 | | | | | |
| | Diethyl phthalate | 15 | 59 | | | | 5,100 | | | | | |
| | Dimethyl phthalate Di-n-butyl phthalate | 15 15 | 59 59 | 3 | 0.016 | 0.013 | 630 | | | | | |
| | Di-n-octyl phthalate | 15 | 59 | 3 | 0.016 | 0.013 | 63 | | 2 | | | |
| | Hexachlorobenzene | 15 | 59 | | | | 0.21 | | 17 | | | |
| | Hexachlorobutadiene | 1 | 1 | | | | 1.2 | | | | | |
| | Hexachlorocyclopentadiene | 15 | 59 | | | | 0.18 | | 17 | | | |
| | Hexachloroethane | 15 | 56 | | | | 1.8 | | 8 | | | |
| | Isophorone | 15 | 59 | 1 | 6.3 | 6.3 | 570 | | | | | |
| | Naphthalene Nitrobenzene | 1 0 | 1 17 | 1 | 0.00071 | 0.00071 | 3.8 5.1 | | 4 | | | |
| | N-Nitrosodimethylamine | 8 7 | 42 | | | | 0.002 | - | 4 42 | | | |
| | N-Nitroso-di-n-propylamine | | 59 | | | | 0.002 | | 17 | | | |
| | N-Nitrosodiphenylamine | 15 | 59 | | | | 110 | | 1 | | | |
| | Pentachlorophenol | 15 | 59 | 3 | 0.0036 | 0.00081 | 1 | | 10 | | | |
| | Phenol | 15 | 59 | 6 | 0.1 | 0.023 | 1,900 | | | | | |
| | 2,4-Dinitrotoluene | 8 | 17 | | | | 1.7 | | 8 | | | |
| | 2,6-Dinitrotoluene | 8 | 17 | 47 | 60 | 0.0000 | 0.36 | ļ | 17 | | | |
| | 2-Methylnaphthalene 1,1,1,2-Tetrachloroethane | 7 15 | 42 59 | 17 | 63 | 0.0006 | 24 | 1 | 2 | | | |
| | 1,1,1,2-Tetrachioroethane | 15 | 59 59 | | | | 810 | | | | | |
| | 1,1,2 - Trichlorotrifluoroethane | 7 | 42 | | 1 | | 4,000 | | | | | |
| | 1,1,2,2-Tetrachloroethane | 15 | 59 | | | | 0.6 | 1 | 2 | | | |
| | 1,1,2-Trichloroethane | 15 | 59 | | | | 0.15 | | 5 | | | |
| | 1,1-Dichloroethane | 15 | 59 | | | | 3.6 | | 1 | | | |
| | 1,1-Dichloroethene | 15 | 57 | | | | 23 | | | | | |
| | 1,1-Dichloropropene | 8 | 17 | | 0.00017 | 0.00010 | 0.0 | ļ | | | | |
| | 1,2,3-Trichlorobenzene | 15 15 | 59 | 6 | 0.00017 | 0.00013 | 6.3 0.0051 | ļ | 2 | | | |
| I | 1,2,3-Trichloropropane | 15 | 59 | | | | T 0.005T | 1 | 11 | | | |

Table 3-10 - Summary of Soil DataBremerton Gas Works Superfund Site
Bremerton, Washington

| Chemical Group | Chemical Constituent | Number of Locations | Number of Samples | Number of Detections | Maximum Detected Concentration (mg/kg) | Minimum Detected Concentration (mg/kg) | Soil PRG (mg/kg) | Detected Concentrations Exceeding the PRG | Number of Non- Detect Results with Reporting Limit Concentrations that Exceed the PRG | Puget Sound Background Metals Concentration (mg/kg)1 | Number of Detected Concentrations Exceeding Puget Sound Background Concentration | Number of Non- Detect Results with Reporting Limit Concentrations that Exceed Puget Sound Background Concentration |
|-----------------------------------|---|------------------------|----------------------|-------------------------|---|---|---------------------|---|---|--|--|--|
| | 1,2,4-Trichlorobenzene | 15 | 59 | 2 | 0.00023 | 0.00014 | 5.8 | | 2 | | | |
| | 1,2,4-Trimethylbenzene | 15 | 59 | 9 | 13.2 | 0.014 | 5.8 | 4 | | | | |
| | 1,2-Dibromo-3-chloropropane | 15 | 59 | | | | 0.0053 | | 19 | | | |
| | 1,2-Dibromoethane (EDB) | 15 | 59 | | | | 0.036 | | 11 | | | |
| | 1,2-Dichlorobenzene 1,2-Dichloroethane (EDC) | 15 15 | 58 59 | | | | 180 0.46 | | 4 | | | |
| | 1,2-Dichloropropane | 15 | 58 | | | | 0.46 | | 2 | | | |
| | 1,3,5-Trimethylbenzene | 15 | 59 | 8 | 5.5 | 0.026 | 78 | | | | | |
| | 1,3-Dichlorobenzene | 9 | 19 | • | 0.0 | 0.020 | | | | | | |
| | 1,3-Dichloropropane | 8 | 17 | | | | 160 | | | | | |
| | 1,4-Dichlorobenzene | 15 | 58 | 1 | 0.00037 | 0.00037 | 2.6 | | 1 | | | |
| | 1,4-Difluorobenzene | 1 | 1 | 1 | 2 | 2 | | | | | | |
| | 2,2-Dichloropropane | 8 | 17 | | | | | | | | | |
| | 2-Butanone | 15 | 59 | 2 | 2.4 | 0.015 | 2,700 | | | | | |
| | 2-Chlorotoluene | 8 | 17 | | | | 160 | | 4 | | | |
| | 2-Hexanone | 15 | 59 | | | | 20 | | 1 | | | |
| | 4-Chlorotoluene | 8 15 | 17 59 | | | | 160 3,300 | | | | | |
| | 4-Methyl-2-pentanone Acetone | 15 | 59 | 30 | 0.064 | 0.0065 | 6,100 | | | | | |
| | Benzene | 15 | 59 | 22 | 12 | 0.00069 | 1.2 | 3 | | | | |
| | Bromobenzene | 8 | 17 | 22 | 12 | 0.00003 | 29 | | | | | |
| | Bromochloromethane | 15 | 59 | | | | 15 | | | | | |
| | Bromodichloromethane | 15 | 59 | | | | 0.29 | | 5 | | | |
| | Bromoform | 15 | 59 | | | | 19 | | | | | |
| | Bromomethane | 15 | 58 | | | | 0.68 | | 2 | | | |
| | Carbon disulfide | 15 | 59 | 4 | 0.0075 | 0.0043 | 77 | | | | | |
| | Carbon tetrachloride | 15 | 59 | | | | 0.65 | | 2 | | | |
| Volatile Organic Compounds (VOCs) | Chlorobenzene | 15 | 59 | 4 | 0 | | 28 | | | | | |
| | Chlorobenzene-d5 Chloroethane | 1 15 | 1 59 | 1 | 2 | 2 | 1,400 | | | | | |
| | Chloroform | 15 | 59 | 3 | 0.044 | 0.00048 | 0.32 | | 5 | | | |
| | Chloromethane | 15 | 59 | 3 | 0.044 | 0.00040 | 11 | | 1 | | | |
| | cis-1,2-Dichloroethene (DCE) | 15 | 59 | | | | 16 | | ' | | | |
| | cis-1,3-Dichloropropene | 15 | 59 | 3 | 0.93 | 0.00063 | | | | | | |
| | Cyclohexane | 7 | 42 | | | | 650 | | | | | |
| | Dibromochloromethane | 15 | 59 | | | | 8.3 | | | | | |
| | Dibromomethane | 8 | 17 | | | | 2.4 | | 1 | | | |
| | Dichlorodifluoromethane | 15 | 59 | | | | 8.7 | | | | | |
| | Ethylbenzene | 15 | 59 | 16 | 24 | 0.00073 | 5.8 | 1 | | | | |
| | Hexachlorobutadiene | 15 | 58 | | | | 1.2 | | 3 | | | |
| | Hexachloroethane Isopropylbenzene | 1 15 | 3 59 | 7 | 1.6 | 0.00094 | 1.8 190 | | | | | |
| | Methyl acetate | 7 | 42 | 1 | 0.16 | 0.00094 | 7,800 | | | | | |
| | Methyl tert-butyl ether (MTBE) | 15 | 59 | ' | 5.10 | 5.10 | 47 | | | | | |
| | Methylcyclohexane | 7 | 42 | 3 | 0.0038 | 0.00037 | · · · · · · | | | | | |
| | Methylene chloride | 15 | 59 | 24 | 1.3 | 0.00058 | 35 | | 1 | | | |
| | n-Butylbenzene | 8 | 17 | 2 | 1.96 | 1.78 | 390 | | | | | |
| | n-Hexane | 8 | 17 | 1 | 0.00121 | 0.00121 | 61 | | | | | |
| | n-Propylbenzene | 8 | 17 | 2 | 0.952 | 0.792 | 380 | | | | | |
| | Pentafluorobenzene | 2 | 3 | 3 | 2 | 0.04 | | | | | | |
| | p-Isopropyltoluene | 8 8 | 17 17 | 4 2 | 1.65 0.915 | 0.493 0.748 | 780 | | | | | |
| | sec-Butylbenzene Styrene | 15 | 59 | 4 | 0.915 | 0.748 | 600 | | | | | |
| | tert-Butylbenzene | 8 | 17 | + | 0.01 | 0.000014 | 780 | | | | | |
| | Tetrachloroethene (PCE) | 15 | 59 | 3 | 0.00059 | 0.00044 | 8.1 | | | | | |
| | Toluene | 15 | 59 | 30 | 7.5 | 0.00026 | 490 | | | | | |
| | trans-1,2-Dichloroethene | 15 | 59 | | | | 160 | | | | | |
| | trans-1,3-Dichloropropene | 15 | 59 | 3 | 0.93 | 0.00063 | | | | | | |
| | Trichloroethene (TCE) | 15 | 59 | 3 | 0.00147 | 0.00044 | 0.41 | | 5 | | | |
| | Trichlorofluoromethane | 15 | 59 | 13 | 0.0078 | 0.0006 | 2300 | | | | | |
| | Vinyl chloride | 15 | 59 | | | | 0.059 | | 11 | | | |
| | m,p-Xylenes | 13 | 50 | 9 | 57 | 0.00052 | | | | | | |

Table 3-10 - Summary of Soil Data

Bremerton Gas Works Superfund Site Bremerton, Washington

| Chemical Group | Chemical Constituent | Number of Locations | | Number of Detections | Maximum Detected Concentration (mg/kg) | Minimum Detected Concentration (mg/kg) | | Detected Concentrations Exceeding the PRG | Number of Non- Detect Results with Reporting Limit Concentrations that Exceed the PRG | Puget Sound Background Metals Concentration (mg/kg)1 | Number of Detected Concentrations Exceeding Puget Sound Background Concentration | Number of Non- Detect Results with Reporting Limit Concentrations that Exceed Puget Sound Background Concentration |
|----------------------------------|----------------------|------------------------|----|-------------------------|---|---|------|---|---|--|--|--|
| | o-Xylene | 13 | 50 | 8 | 55 | 0.00049 | 65 | | | | | |
| | Xylenes (total) | 8 | 17 | 7 | 16.7 | 0.353 | 58 | | | | | |
| | Naphthalene | 14 | 49 | 33 | 708 | 0.00059 | 3.8 | 7 | | | | |
| | Aroclor 1016 | 8 | 17 | | | | 0.41 | | | | | |
| | Aroclor 1221 | 8 | 17 | | | | 0.2 | | | | | |
| | Aroclor 1232 | 8 | 17 | | | | 0.17 | | | | | |
| | Aroclor 1242 | 8 | 17 | | | | 0.23 | | | | | |
| Polychlorinated Biphenyls (PCBs) | Aroclor 1248 | 8 | 17 | | | | 0.23 | | | | | |
| | Aroclor 1254 | 8 | 17 | | | | 0.12 | | | | | |
| | Aroclor 1260 | 8 | 17 | | | | 0.24 | | | | | |
| | Aroclor 1262 | 8 | 17 | | | | | | | | | |
| | Aroclor 1268 | 8 | 17 | | | | | | | | | |

Reference: U.S. Environmental Protection Agency (EPA), 1993, Provisional Guidance for Quantitative Risk Assessment of Polycyclic Aromatic Hydrocarbons, Office of Research and Development, Office of Health and Environmental Assessment, Washington, DC, EPA/600/R-93/089.

Table 3-11 - Summary of Groundwater DataBremerton Gas Works Superfund Site Bremerton, Washington

| | | | | | Maximum Detected Concentration | | Groundwater | Number of Detected Concentrations Exceeding the | Number of Non- Detect Results with Reporting Limit Concentrations that Exceed the Groundwater | Surface Water | | Concentrations that Exceed the Surface |
|---|--------------------------------|-----------|---------|------------|--------------------------------------|--------|-------------|---|---|---------------|-------------------|---|
| Chemical Group | Chemical Constituent | Locations | Samples | Detections | (ug/L) | (ug/L) | PRG (ug/L) | Groundwater PRG | PRG | PRG (ug/L) | Surface Water PRG | Water PRG |
| | Gasoline-Range Hydrocarbons | 10 | 10 | 7 | 10,600 | 63.5 | | | | | | |
| Total Petroleum Hydrocarbons (TPH) | Diesel-Range Hydrocarbons | 11 | 11 | 6 | 18,500 | 170 | | | | | | |
| | Oil-Range Hydrocarbons | 11 | 11 | 1 | 160 | 160 | | | | | | |
| | Antimony | 10 | 10 | 2 | 0.4 | 0.3 | 0.78 | | 8 | 90 | | |
| | Arsenic | 10 | 10 | 10 | 26 | 0.6 | 0.052 | 10 | | 0.14 | 10 | |
| | Barium | 2 | 2 | 2 | 173 | 35.7 | 380 | | | | | |
| | Beryllium | 10 | 10 | 3 | 1.08 | 0.37 | 2.5 | | | 0.66 | 2 | 7 |
| | Cadmium | 10 | 10 | 2 | 0.16 | 0.05 | 0.92 | | 8 | 8.8 | | |
| | Chromium (Total) | 10 | 10 | 10 | 228 | 1.34 | 100 | 2 | | 42 | 3 | |
| | Chromium (VI) | 8 | 8 | 7 | 90 | 6 | 0.035 | 7 | 1 | 50 | 2 | |
| | Cobalt | 2 | 2 | 2 | 8.3 | 1.4 | 0.6 | 2 | | | | |
| Metals | Copper | 10 | 10 | 10 | 143 | 1.05 | 80 | 2 | | 3.1 | 8 | |
| Metals | Lead | 10 | 10 | 8 | 21.6 | 0.44 | 15 | 2 | | 8.1 | 2 | |
| | Manganese | 2 | 2 | 2 | 3,020 | 98.1 | 43 | 2 | | | | |
| | Mercury | 8 | 8 | 1 | 0.246 | 0.246 | 0.063 | 1 | 7 | 0.94 | | |
| | Nickel | 10 | 10 | 10 | 232 | 1.65 | 39 | 3 | | 8.2 | 7 | |
| | Selenium | 10 | 10 | 1 | 3.64 | 3.64 | 10 | | | 71 | | |
| | Silver | 10 | 10 | 1 | 0.07 | 0.07 | 9.4 | | | 1.9 | | |
| | Thallium | 10 | 10 | 1 | 0.26 | 0.26 | 0.02 | 1 | 9 | 6.3 | | |
| | Vanadium | 2 | 2 | 2 | 78.2 | 3.7 | 8.6 | 1 | | | | |
| | Zinc | 10 | 10 | 8 | 185 | 4.5 | 600 | | | 81 | 2 | |
| TCLP Metals | Mercury | 8 | 8 | 1 | 0.246 | 0.246 | 0.063 | 1 | 7 | 0.94 | | |
| | Acenaphthene | 9 | 9 | 5 | 485 | 1.1 | 53 | 1 | | 30 | 1 | |
| | Acenaphthylene | 10 | 10 | 6 | 34.9 | 0.222 | | | | 4,840 | | |
| | Anthracene | 10 | 10 | 5 | 120 | 0.4 | 180 | | | 100 | 1 | |
| | Benzo(g,h,i)perylene | 10 | 10 | 5 | 25.6 | 0.0979 | | | | 7.64 | 1 | |
| | Dibenzofuran | 10 | 10 | 2 | 31.8 | 0.29 | 0.79 | 1 | 7 | 4 | 1 | 7 |
| Delvevelie Aremetie Hydrocerbone (DAHe) | Fluoranthene | 10 | 10 | 6 | 122 | 0.26 | 80 | 1 | | 6 | 2 | |
| Polycyclic Aromatic Hydrocarbons (PAHs) | Fluorene | 10 | 10 | 7 | 184 | 0.102 | 29 | 1 | | 10 | 2 | |
| | Phenanthrene | 10 | 10 | 5 | 377 | 1.04 | | | | 1.5 | 3 | |
| | Pyrene | 10 | 10 | 7 | 34.5 | 0.174 | 12 | 2 | | 8 | 2 | |
| | 1-Methylnaphthalene | 8 | 8 | 4 | 970 | 0.813 | 1.1 | 3 | 11 | 2.1 | 3 | 11 |
| | 2-Methylnaphthalene | 10 | 10 | 6 | 1,430 | 0.13 | 3.6 | 1 | | 4.2 | 1 | |
| | Naphthalene | 2 | 2 | | | | 0.17 | | | 1.4 | | |
| | Benz(a)anthracene | 10 | 10 | 6 | 39.3 | 0.0168 | 0.012 | 6 | 2 | 0.00016 | 6 | 4 |
| | Benzo(a)pyrene | 10 | 10 | 6 | 37.6 | 0.0247 | 0.0034 | 6 | 4 | 0.000016 | 6 | 4 |
| | Benzo(b)fluoranthene | 10 | 10 | 4 | 0.657 | 0.0968 | 0.034 | 4 | 3 | 0.00016 | 4 | 6 |
| | Benzo(k)fluoranthene | 10 | 10 | 5 | 0.615 | 0.0602 | 0.34 | 2 | 1 | 0.0016 | 5 | 5 |
| Carcinogenic PAHs (cPAHs) | Chrysene | 10 | 10 | 6 | 40.8 | 0.0372 | 3.4 | 1 | | 0.016 | 6 | 2 |
| | Dibenzo(a,h)anthracene | 10 | 10 | 4 | 0.189 | 0.0437 | 0.0034 | 4 | 6 | 0.000016 | 4 | 6 |
| | Indeno(1,2,3-cd)pyrene | 10 | 10 | 4 | 0.467 | 0.0874 | 0.034 | 4 | 3 | 0.00016 | 4 | 6 |
| | Total cPAHs TEQ (ND = 0) | 10 | 10 | 6 | 41.9 | 0.0328 | 0.0034 | 6 | | 0.000016 | 6 | |
| | Total cPAHs TEQ (ND = 1/2 RDL) | 10 | 10 | 6 | 43.8 | 0.0342 | 0.0034 | 6 | | 0.000016 | 6 | |

Table 3-11 - Summary of Groundwater DataBremerton Gas Works Superfund Site Bremerton, Washington

| Chemical Group | Chemical Constituent | Number of Locations | | Number of Detections | | Minimum Detected Concentration (ug/L) | Groundwater PRG (ug/L) | Number of Detected Concentrations Exceeding the Groundwater PRG | Number of Non- Detect Results with Reporting Limit Concentrations that Exceed the Groundwater PRG | Surface Water PRG (ug/L) | Number of Detected Concentrations Exceeding the Surface Water PRG | Concentrations that Exceed the Surface |
|---------------------------------------|-----------------------------------|------------------------|----|-------------------------|------|--|---------------------------|--|--|-----------------------------|--|--|
| | 1,1'-Biphenyl | 2 | 2 | | | | 0.083 | | 2 | 14 | | |
| | 1,2,4,5-Tetrachlorobenzene | 2 | 2 | | | | 0.17 | | 2 | 1.1 | | |
| | 1,2,4-Trimethylbenzene | 1 | 1 | | | | 1.5 | | | 19 | | |
| | 2,3,4,6-Tetrachlorophenol | 2 | 2 | | | | 24 | | | 1.2 | | |
| | 2,4,5-Trichlorophenol | 10 | 10 | | | | 120 | | | 12 | | |
| | 2,4,6-Trichlorophenol | 10 | 10 | | | | 1.2 | | 8 | 0.28 | | 10 |
| | 2,4-Dichlorophenol | 10 | 10 | | | | 4.6 | | 8 | 10 | | 1 |
| | 2,4-Dimethylphenol | 10 | 10 | | | | 36 | | | 97 | | |
| | 2,4-Dinitrophenol | 10 | 10 | | | | 3.9 | | 8 | 100 | | |
| | 2-Chloronaphthalene | 10 | 10 | | | | 75 | | | 100 | | |
| | 2-Chlorophenol | 10 | 10 | | | | 9.1 | | 8 | 17 | | |
| | 2-Methylphenol | 8 | 8 | | | | 93 | | | 67 | | |
| | 2-Nitroaniline | 10 | 10 | | | | 19 | | | | | |
| | 2-Nitrophenol | 10 | 10 | | | | | | | 2,940 | | |
| | 3 & 4 Methylphenol | 8 | 8 | | | | | | | | | |
| | 3,3'-Dichlorobenzidine | 10 | 10 | | | | 0.13 | | 10 | 0.0033 | | 10 |
| | 3-Nitroaniline | 10 | 10 | | | | | | | | | |
| | 4,6-Dinitro-2-methylphenol | 10 | 10 | | | | 0.15 | | 10 | 7 | | 8 |
| | 4-Bromophenyl phenyl ether | 10 | 10 | | | | | | | 1.5 | | 8 |
| | 4-Chloro-3-methylphenol | 10 | 10 | | | | 140 | | | 36 | | |
| | 4-Chloroaniline | 10 | 10 | | | | 0.37 | | 10 | 232 | | |
| | 4-Chlorophenyl phenyl ether | 10 | 10 | | | | | | | | | |
| | 4-Methylphenol | 2 | 2 | | | | 190 | | | 25 | | |
| | 4-Nitroaniline | 10 | 10 | | | | 3.8 | | 8 | | | |
| | 4-Nitrophenol | 10 | 10 | | | | | | | 60 | | |
| | Acenaphthene | 1 | 1 | 1 | 4.9 | 4.9 | 53 | | | 30 | 1 | |
| | Acetophenone | 2 | 2 | | | | 190 | | | | | |
| Other Semivolaitile Organic Compounds | Aniline | 8 | 8 | | | | 13 | | | 2.2 | | 8 |
| (SVOCs) | Atrazine | 2 | 2 | | | | 0.3 | | 2 | 1.8 | | |
| | Benzaldehyde | 2 | 2 | | | | 19 | | | | | |
| | Benzidine | 2 | 2 | | | | 0.00011 | | 2 | 0.000023 | | 2 |
| | Benzoic acid | 8 | 8 | | | | 7,500 | | | 42 | | |
| | Benzyl alcohol | 8 | 8 | | | | 200 | | | 8.6 | | 8 |
| | Benzyl butyl phthalate | 10 | 10 | 1 | 0.33 | 0.33 | 16 | | | 0 | 1 | 9 |
| | Bis(2-chloro-1-methylethyl) ether | 10 | 10 | | | | 71 | | | 900 | | |
| | Bis(2-chloroethoxy)methane | 10 | 10 | | | | 5.9 | | 8 | | | |
| | Bis(2-chloroethyl) ether | 10 | 10 | | | | 0.014 | | 10 | 0.06 | | 10 |
| | Bis(2-ethylhexyl) phthalate | 10 | 10 | 2 | 0.5 | 0.33 | 5.6 | | 8 | 0.046 | 2 | 8 |
| | Caprolactam | 2 | 2 | 1 | 0.71 | 0.71 | 990 | | | | | |
| | Carbazole | 10 | 10 | 1 | 1.3 | 1.3 | | | | | | |
| | Diethyl phthalate | 10 | 10 | | | | 1,500 | | | 200 | | |
| | Dimethyl phthalate | 10 | 10 | | | | | | | 600 | | |
| | Di-n-butyl phthalate | 10 | 10 | | | | 90 | | | 8 | | 8 |
| | Di-n-octyl phthalate | 10 | 10 | | | | 20 | | | 22 | | |
| | Hexachlorobenzene | 10 | 10 | | | | 0.0098 | | 10 | 0.000005 | | 10 |
| | Hexachlorocyclopentadiene | 10 | 10 | | | | 0.041 | | 10 | 1 | | 8 |
| | Hexachloroethane | 10 | 10 | | | | 0.33 | | 10 | 0.02 | | 10 |
| | Isophorone | 10 | 10 | | | | 78 | | | 110 | | |
| | Nitrobenzene | 8 | 8 | | | | 0.14 | | 8 | 100 | | |
| | N-Nitrosodimethylamine | 2 | 2 | | | | 0.00011 | | 2 | 0 | | 2 |
| | N-Nitroso-di-n-propylamine | 10 | 10 | | | | 0.011 | | 10 | 0.058 | | 10 |

Table 3-11 - Summary of Groundwater DataBremerton Gas Works Superfund Site Bremerton, Washington

| Chemical Group | Chemical Constituent | Number of Locations | | Number of Detections | | Minimum Detected Concentration (ug/L) | Groundwater PRG (ug/L) | Number of Detected Concentrations Exceeding the Groundwater PRG | Number of Non- Detect Results with Reporting Limit Concentrations that Exceed the Groundwater PRG | Surface Water PRG (ug/L) | Number of Detected Concentrations Exceeding the Surface Water PRG | Concentrations that Exceed the Surface |
|-----------------------------------|----------------------------------|---------------------|----|-------------------------|--|--|---------------------------|--|--|-----------------------------|--|---|
| | N-Nitrosodiphenylamine | 10 | 10 | | | | 12 | | | 0.69 | | 8 |
| | Pentachlorophenol | 10 | 10 | 2 | 11.4 | 0.1 | 0.041 | 2 | 8 | 0.002 | 2 | 8 |
| | Phenol | 10 | 10 | 3 | 81.6 | 75.5 | 580 | | | 70,000 | | |
| | 2,4-Dinitrotoluene | 8 | 8 | | | | 0.24 | | 8 | 0.18 | | 8 |
| | 2,6-Dinitrotoluene | 8 | 8 | | | | 0.049 | | 8 | 81 | | |
| | 1,1,1,2-Tetrachloroethane | 10 | 10 | | | | 0.57 | | | | | |
| | 1,1,1-Trichloroethane | 10 | 10 | | | | 200 | | | 50000 | | |
| | 1,1,2 - Trichlorotrifluoroethane | 2 | 2 | | | | 5,500 | | | | | |
| | 1,1,2,2-Tetrachloroethane | 10 | 10 | | | | 0.076 | | 10 | 0.3 | | 8 |
| | 1,1,2-Trichloroethane | 10 | 10 | | | | 0.041 | | 10 | 0.9 | | † |
| | 1,1-Dichloroethane | 10 | 10 | | | | 2.8 | | ' | 47 | | |
| | 1,1-Dichloroethene | 9 | 9 | | 1 | | 7 | | | 4,000 | | |
| | 1,1-Dichloropropene | 8 | 8 | | | | 1 | | | 4,000 | | |
| | 1,2,3-Trichlorobenzene | 10 | 10 | | | | 0.7 | | 8 | 8 | | |
| | 1,2,3-Trichloropenzene | 10 | 10 | | | | 0.00075 | | 10 | • | | |
| | | | | | | | | | | 0.037 | | 10 |
| - | 1,2,4-Trichlorobenzene | 10 | 10 | - | 170 | 2.52 | 0.4 | <u> </u> | 10 | | | 10 |
| | 1,2,4-Trimethylbenzene | 9 | 9 | 5 | 179 | 3.52 | 1.5 | 5 | 40 | 19 | | |
| | 1,2-Dibromo-3-chloropropane | 10 | 10 | | | | 0.00033 | | 10 | | | |
| - | 1,2-Dibromoethane (EDB) | 10 | 10 | | | | 0.0075 | | 10 | 200 | | |
| - | 1,2-Dichlorobenzene | 10 | 10 | | . == | | 30 | | _ | 800 | | |
| - | 1,2-Dichloroethane (EDC) | 10 | 10 | 3 | 4.72 | 0.93 | 0.17 | 3 | 7 | 73 | | |
| - | 1,2-Dichloropropane | 10 | 10 | | | | 0.44 | | | 3.1 | | |
| _ | 1,3,5-Trimethylbenzene | 10 | 10 | 5 | 30 | 0.53 | 12 | 1 | | 71 | | |
| _ | 1,3-Dichlorobenzene | 10 | 10 | | | | | | | 2 | | |
| | 1,3-Dichloropropane | 8 | 8 | | | | 37 | | | | | |
| _ | 1,4-Dichlorobenzene | 10 | 10 | | | | 0.48 | | | 200 | | |
| | 2,2-Dichloropropane | 8 | 8 | | | | | | | | | |
| | 2-Butanone | 10 | 10 | | | | 560 | | | 2,200 | | |
| | 2-Chlorotoluene | 8 | 8 | | | | 24 | | | | | |
| | 2-Hexanone | 10 | 10 | | | | 3.8 | | 2 | 99 | | |
| | 4-Chlorotoluene | 8 | 8 | | | | 25 | | | | | |
| | 4-Methyl-2-pentanone | 10 | 10 | | | | 630 | | | 170 | | |
| | Acetone | 10 | 10 | | | | 1,400 | | | 1,700 | | |
| | Benzene | 10 | 10 | 8 | 950 | 2.23 | 0.46 | 8 | | 1.6 | 8 | |
| | Bromobenzene | 8 | 8 | | | | 6.2 | | | | | |
| | Bromochloromethane | 10 | 10 | | | | 8.3 | | | | | |
| Γ | Bromodichloromethane | 10 | 10 | | | | 0.13 | | 10 | 2.8 | | |
| Γ | Bromoform | 10 | 10 | | | | 3.3 | | | 12 | | |
| Γ | Bromomethane | 10 | 10 | | | | 0.75 | | 8 | 2,400 | | |
| Γ | Carbon disulfide | 10 | 10 | | | | 81 | | | 0.92 | | |
| Volatile Organic Compounds (VOCs) | Carbon tetrachloride | 10 | 10 | 1 | 0.66 | 0.66 | 0.46 | 1 | | 0.35 | 1 | |
| | Chlorobenzene | 10 | 10 | | | | 7.8 | | | 200 | | |
| ļ | Chloroethane | 10 | 10 | | | | 2,100 | | | | | |
| ļ | Chloroform | 10 | 10 | 3 | 2.84 | 0.2 | 0.22 | 2 | 2 | 600 | | |
| | Chloromethane | 10 | 10 | | | | 19 | | | 2,700 | | |
| | cis-1,2-Dichloroethene (DCE) | 10 | 10 | 3 | 1.29 | 0.37 | 3.6 | | | | | |
| | cis-1,3-Dichloropropene | 10 | 10 | <u> </u> | 1 | | | | | 1.2 | | |
| | Cyclohexane | 2 | 2 | 1 | 0.38 | 0.38 | 1,300 | | | <u>-</u> | | 1 |
| | Dibromochloromethane | 10 | 10 | | 1 | | 0.87 | | | 2.2 | | 1 |
| | Dibromomethane | 8 | 8 | | 1 | | 0.83 | | | | | <u> </u> |
| | | <u> </u> | 10 | | | | 20 | + | | | ļ | + |

Table 3-11 - Summary of Groundwater Data

Bremerton Gas Works Superfund Site Bremerton, Washington

| Chemical Group | Chemical Constituent | Number of Locations | | Number of Detections | Maximum Detected Concentration (ug/L) | Minimum Detected Concentration (ug/L) | Groundwater PRG (ug/L) | Number of Detected Concentrations Exceeding the Groundwater PRG | Number of Non- Detect Results with Reporting Limit Concentrations that Exceed the Groundwater PRG | Surface Water PRG (ug/L) | Number of Detected Concentrations Exceeding the Surface Water PRG | Number of Non- Detect Results with Reporting Limit Concentrations that Exceed the Surface Water PRG |
|----------------------------------|--------------------------------|------------------------|----|----------------------|--|--|---------------------------|--|--|-----------------------------|--|--|
| | Ethylbenzene | 10 | 10 | 7 | 322 | 0.53 | 1.5 | 6 | | 31 | 4 | |
| | Hexachlorobutadiene | 10 | 10 | | | | 0.14 | | 10 | 0.01 | 1 | 10 |
| | Isopropylbenzene | 10 | 10 | 6 | 37.4 | 3 | 45 | | | 2.6 | 6 | |
| | Methyl acetate | 2 | 2 | | | | 2,000 | | | | | |
| | Methyl tert-butyl ether (MTBE) | 10 | 10 | | | | 14 | | | 11,070 | | |
| | Methylcyclohexane | 2 | 2 | | | | | | | | | |
| | Methylene chloride | 10 | 10 | | | | 5 | | | 100 | | |
| | n-Butylbenzene | 8 | 8 | 4 | 5.3 | 0.48 | 100 | | | | | |
| | n-Hexane | 8 | 8 | 1 | 1.17 | 1.17 | 150 | | | 0.58 | 1 | 7 |
| | n-Propylbenzene | 8 | 8 | 4 | 9.2 | 2.38 | 66 | | | 128 | | |
| | p-Isopropyltoluene | 8 | 8 | 4 | 8.44 | 0.27 | | | | 85 | | |
| | sec-Butylbenzene | 8 | 8 | 5 | 4.43 | 0.32 | 200 | | | | | |
| | Styrene | 10 | 10 | | | | 100 | | | 32 | | |
| | tert-Butylbenzene | 8 | 8 | | | | 69 | | | | | |
| | Tetrachloroethene (PCE) | 10 | 10 | | | | 4.1 | | | 2.9 | | |
| | Toluene | 10 | 10 | 6 | 41.9 | 0.45 | 110 | | | 130 | | |
| | trans-1,2-Dichloroethene | 10 | 10 | | | | 36 | | | 1,000 | | |
| | trans-1,3-Dichloropropene | 10 | 10 | | | | | | | 1.2 | | |
| | Trichloroethene (TCE) | 10 | 10 | 6 | 4.79 | 0.33 | 0.28 | 6 | | 0.7 | 1 | |
| | Trichlorofluoromethane | 10 | 10 | | | | 520 | | | | | |
| | Vinyl chloride | 10 | 10 | | | | 0.019 | | 10 | 0.18 | | 10 |
| | m,p-Xylenes | 10 | 10 | 6 | 383 | 0.74 | | | | | | |
| | o-Xylene | 10 | 10 | 6 | 211 | 4.91 | 19 | 2 | | | | |
| | Xylenes (total) | 8 | 8 | 5 | 593 | 8.29 | 19 | 4 | | 19 | 4 | |
| | Naphthalene | 8 | 8 | 8 | 5,270 | 0.47 | 0.17 | 8 | | 1.4 | | |
| | Aroclor 1016 | 8 | 8 | | | | 0.14 | | | | | |
| | Aroclor 1221 | 8 | 8 | | | | 0.0047 | | 8 | | | |
| | Aroclor 1232 | 8 | 8 | | | | 0.0047 | | 8 | | | |
| | Aroclor 1242 | 8 | 8 | | | | 0.0078 | | 8 | | | |
| Polychlorinated Biphenyls (PCBs) | Aroclor 1248 | 8 | 8 | | | | 0.0078 | | 8 | | | |
| | Aroclor 1254 | 8 | 8 | | | | 0.0078 | | 8 | | | |
| | Aroclor 1260 | 8 | 8 | | | | 0.0078 | | 8 | | | |
| | Aroclor 1262 | 8 | 8 | | | | | | | | | |
| | Aroclor 1268 | 8 | 8 | | | | | | | | | |

Reference: U.S. Environmental Protection Agency (EPA), 1993, Provisional Guidance for Quantitative Risk Assessment of Polycyclic Aromatic Hydrocarbons, Office of Research and Development, Office of Health and Environmental Assessment, Washington, DC, EPA/600/R-93/089.

Table 3-12 - Summary of Sediment DataBremerton Gas Works Superfund Site

| | | Number of | Number of | Number of | Maximum Detected Concentration | Minimum Detected Concentration | Sediment PRG | Puget Sound Background Sediment Concentration ¹ | Number of Detected Concentrations Exceeding the | Number of Detected Concentrations Exceeding Puget Sound Background Metals |
|--|--------------------------------|-----------|-----------|------------|--------------------------------------|--------------------------------------|-----------------|---|--|--|
| Chemical Group | Chemical Constituent | Locations | Samples | Detections | (ug/kg) | (ug/kg) | (ug/kg) | (ug/kg) | PRG | Concentration |
| | Gasoline Range Hydrocarbons | 5 | 5 | 0 | NA | NA | | | | |
| TPH | Diesel Range Hydrocarbons | 5 | 5 | 4 | 240000 | 63000 | | | | |
| | Oil Range Hydrocarbons | 5 | 5 | 5 | 620000 | 21000 | | | | |
| | Aluminum | 5 | 5 | 5 | 9030000 | 6020000 | | | | |
| | Antimony | 1 | 1 | 1 | 3900 | 3900 | 2000 | 5000 | 1 | |
| | Arsenic | 5 | 5 | 5 | 5100 | 1500 | 57000 | 11000 | | |
| | Barium | 5 | 5 | 5 | 47000 | 13300 | | | | |
| L | Beryllium | 5 | 5 | 5 | 2700 | 1900 | | | | |
| | Cadmium | 5 | 5 | 0 | NA | NA | 5100 | 800 | | |
| | Calcium | 5 | 5 | 5 | 33600000 | 2390000 | | | | |
| | Chromium (Total) | 5 | 5 | 5 | 21200 | 16600 | 260000 | 62000 | | |
| | Cobalt | 5 | 5 | 5 | 26300 | 3000 | 50000 | 11000 | | |
| | Copper | 5 | 5 | 5 | 71700 | 8600 | 390000 | 45000 | | |
| | Iron | 5 | 5 | 5 | 15900000 | 9730000 | 20000000 | | | |
| Metals | Lead | 5 | 5 | 5 | 30000 | 8900 | 450000 | 21000 | | |
| | Magnesium | 5 | 5 | 5 | 4640000 | 3350000 | | | | |
| | Manganese | 5 | 5 | 5 | 180000 | 135000 | 460000 | | | |
| | Mercury | 3 | 3 | 3 | 100 | 27.8 | 410 | 200 | | |
| | Nickel | 5 | 5 | 5 | 52600 | 21400 | 20900 | 50000 | 5 | 1 |
| Ī | Potassium | 5 | 5 | 5 | 603000 | 415000 | | | | |
| Ī | Selenium | 5 | 5 | 1 | 400 | 400 | 2000 | 780 | | |
| | Silver | 5 | 5 | 0 | NA | NA | 6100 | 240 | | |
| T I | Sodium | 5 | 5 | 5 | 1930000 | 605000 | | | | |
| F | Thallium | 5 | 5 | 0 | NA | NA | | | | |
| The state of the s | Vanadium | 5 | 5 | 5 | 36500 | 21600 | | 45000 | | |
| ŀ | Zinc | 5 | 5 | 5 | 79900 | 23200 | 410000 | 93000 | | |
| | Acenaphthene | 48 | 63 | 61 | 160000 | 0.4 | 500 | | 16 | |
| <u> </u> | Acenaphthylene | 51 | 66 | 66 | 840000 | 0.7 | 1300 | | 33 | |
| F | Anthracene | 51 | 66 | 66 | 680000 | 0.3 | 960 | | 41 | |
| ŀ | Benzo(g,h,i)perylene | 51 | 66 | 66 | 260000 | 0.9 | 670 | | 50 | |
| ŀ | Dibenzofuran | 5 | 5 | 4 | 74 | 58 | 540 | | | |
| ŀ | Fluoranthene | 46 | 61 | 61 | 1100000 | 1.6 | 1700 | | 45 | |
| ŀ | Fluorene | 51 | 66 | 65 | 600000 | 0.3 | 540 | | 36 | |
| ŀ | Phenanthrene | 51 | 66 | 66 | 1700000 | 2.6 | 1500 | | 46 | |
| H | Pyrene | 51 | 66 | 66 | 1400000 | 1.6 | 2600 | | 48 | |
| F | 2-Methylnaphthalene | 5 | 5 | 5 | 1200 | 19 | 670 | | 1 | |
| F | Naphthalene | 46 | 61 | 61 | 1700000 | 5.4 | 2100 | | 23 | |
| PAHs | Benz(a)anthracene | 51 | 66 | 66 | 310000 | 0.3 | 1300 | | 46 | |
| PARS | Benzo(a)pyrene | 51 | 66 | 66 | 400000 | 0.5 | 1600 | | 47 | |
| } | Benzo(b)fluoranthene | 51 | 66 | 66 | 200000 | 0.4 | 10400 | | 17 | |
| } | Benzo(k)fluoranthene | 51 | 66 | 65 | 93000 | 0.4 | 240 | | 50 | |
| } | * * | 51 | 66 | 66 | 270000 | 0.5 | 1400 | | 47 | |
| | Chrysene | 51 | 66 | 65 | | | 230 | | 47 | |
| | Dibenzo(a,h)anthracene | | | | 38000 | 0.2 | | | | |
| | Indeno(1,2,3-cd)pyrene | 51 | 66 | 66 | 190000 | 0.4 | 600 | | 49 | |
| | Total cPAHs TEQ (ND = 0) | 51 | 66 | 66 | 509200 | 0.6 | 1600 | 21 | 49 | |
| Ļ | Total cPAHs TEQ (ND = 1/2 RDL) | 51 | 66 | 66 | 509200 | 0.9 | 1600 | 21 | 49 | |
| Ļ | Total HPAHs | 46 | 61 | 61 | 4361000 | 6.2 | 12000 | | 45 | |
| Į | Total LPAHs | 46 | 61 | 61 | 5596000 | 10.1 | 5200 | | 39 | |
| | Total PAHs | 46 | 61 | 61 | 8890000 | 16.3 | 4022 | | 48 | I |

Table 3-12 - Summary of Sediment DataBremerton Gas Works Superfund Site

| Chemical Group | Chemical Constituent | Number of Locations | Number of Samples | Number of Detections | Maximum Detected Concentration (ug/kg) | Minimum Detected Concentration (ug/kg) | Sediment PRG (ug/kg) | Puget Sound Background Sediment Concentration ¹ (ug/kg) | Number of Detected Concentrations Exceeding the PRG | Number of Detected Concentrations Exceeding Puget Sound Background Metals Concentration |
|----------------|-----------------------------------|------------------------|----------------------|-------------------------|---|---|----------------------------|--|---|---|
| chemical Group | | | | | 1 5 5 | | , | (48/ 48/ | 1 1.10 | Concentration |
| - | 1,1'-Biphenyl | 5 5 | 5 | 4 0 | 110 NA | 60 NA | 1220 47000 | | | |
| - | 1,2,4,5-Tetrachlorobenzene | 8 | 9 | 0 | NA NA | NA NA | 31 | | - | |
| - | 1,2,4-Trichlorobenzene | | 9 | | | | | | | |
| - | 1,2-Dichlorobenzene | 8 1 | 1 | 0 1 | NA 34 | NA 21 | 35 | | | |
| - | 1,3,5-Trimethylbenzene | 5 | 5 | 0 | 21 NA | NA | 0.40 | | | |
| - | 1,3-Dichlorobenzene | 2 | 2 | 2 | | | 842 | | - | |
| - | 1,4-Dichlorobenzene | | | | 23 | 22 | 110 | | | |
| | 1,4-Dioxane | 5 | 5 | 0 | NA NA | NA NA | 119 | | | |
| | 2,3,4,6-Tetrachlorophenol | 5 | 5 | | NA | NA | 284 | | | |
| - | 2,4,5-Trichlorophenol | 5 | 5 | 0 | NA | NA | 819 | | | |
| - | 2,4,6-Trichlorophenol | 5 | 5 | 0 | NA | NA | 2650 | | | |
| - | 2,4-Dichlorophenol | 5 | 5 | 0 | NA | NA | 117 | | | |
| - | 2,4-Dimethylphenol | 5 | 5 | 0 | NA | NA | 29 | | | |
| | 2,4-Dinitrophenol | 5 | 5 | 0 | NA | NA | 6.21 | | | |
| | 2-Chloronaphthalene | 5 | 5 | 0 | NA | NA | 417 | | | |
| | 2-Chlorophenol | 5 | 5 | 0 | NA | NA | 344 | | | |
| | 2-Nitroaniline | 5 | 5 | 0 | NA | NA | | | | |
| | 2-Nitrophenol | 5 | 5 | 0 | NA | NA | | | | |
| | 3,3'-Dichlorobenzidine | 5 | 5 | 0 | NA | NA | 2060 | | | |
| | 3-Nitroaniline | 5 | 5 | 0 | NA | NA | | | | |
| | 4,6-Dinitro-2-methylphenol | 5 | 5 | 0 | NA | NA | 104 | | | |
| | 4-Bromophenyl phenyl ether | 5 | 5 | 0 | NA | NA | 1230 | | | |
| | 4-Chloro-3-methylphenol | 5 | 5 | 0 | NA | NA | 388 | | | |
| | 4-Chloroaniline | 5 | 5 | 0 | NA | NA | 146 | | | |
| Other | 4-Chlorophenyl phenyl ether | 5 | 5 | 0 | NA | NA | | | | |
| SVOCs | 4-Methylphenol | 5 | 5 | 2 | 17 | 17 | 670 | | | |
| | 4-Nitroaniline | 5 | 5 | 0 | NA | NA | | | | |
| | 4-Nitrophenol | 5 | 5 | 0 | NA | NA | 13.3 | | | |
| | Acenaphthene | 48 | 63 | 61 | 160000 | 0.4 | 500 | | 16 | |
| | Acetophenone | 5 | 5 | 0 | NA | NA | | | | |
| | Atrazine | 5 | 5 | 0 | NA | NA | 6.62 | | | |
| | Benzaldehyde | 5 | 5 | 2 | 38 | 19 | | | | |
| | Benzidine | 5 | 5 | 0 | NA | NA | | | | |
| | Benzyl butyl phthalate | 5 | 5 | 0 | NA | NA | 63 | | | |
| | Bis(2-chloro-1-methylethyl) ether | 5 | 5 | 0 | NA | NA | | | | |
| | Bis (2-chloroethoxy) methane | 5 | 5 | 0 | NA | NA | | | | |
| | Bis(2-chloroethyl) ether | 5 | 5 | 0 | NA | NA | 3520 | | | |
| | Bis(2-ethylhexyl) phthalate | 5 | 5 | 1 | 42 | 42 | 1300 | | | |
| | Caprolactam | 5 | 5 | 0 | NA | NA | | | | |
| | Carbazole | 5 | 5 | 4 | 110 | 69 | | | | |
| | Dibenzofuran | 5 | 5 | 4 | 74 | 58 | 540 | | | |
| | Diethyl phthalate | 5 | 5 | 0 | NA | NA | 200 | | | |
| ļ | Dimethyl phthalate | 5 | 5 | 0 | NA | NA | 71 | | | |
| | Di-n-butyl phthalate | 5 | 5 | 0 | NA | NA | 1400 | | | |
| | Di-n-octyl phthalate | 5 | 5 | 0 | NA | NA | 6200 | | | |
| | Hexachlorobenzene | 5 | 5 | 0 | NA | NA | 22 | | | |
| ļ | Hexachlorobutadiene | 8 | 9 | 0 | NA | NA | 11 | | | |
| ŀ | Hexachlorocyclopentadiene | 5 | 5 | 0 | NA NA | NA | 139 | | 1 | |
| | Hexachloroethane | 3 | 3 | 0 | NA NA | NA NA | 804 | | | |
| | Isophorone | 5 | 5 | 0 | NA NA | NA NA | 432 | | | |
| } | Naphthalene | 46 | 61 | 61 | 1700000 | 5.4 | 2100 | | 23 | |

Table 3-12 - Summary of Sediment DataBremerton Gas Works Superfund Site

| Chemical Group | Chemical Constituent | Number of Locations | Number of Samples | Number of Detections | Maximum Detected Concentration (ug/kg) | Minimum Detected Concentration (ug/kg) | Sediment PRG (ug/kg) | Puget Sound Background Sediment Concentration ¹ (ug/kg) | Number of Detected Concentrations Exceeding the PRG | Number of Detected Concentrations Exceeding Puget Sound Background Metals Concentration |
|----------------|----------------------------------|------------------------|----------------------|-------------------------|---|---|----------------------------|--|---|---|
| chemical Group | | <u> </u> | | | | | (48/ 48/ | (48/18/ | 1 110 | Concentration |
| Other | N-Nitrosodimethylamine | 5 | 5 | 0 | NA NA | NA NA | | | | |
| SVOCs | N-Nitroso-di-n-propylamine | 5 | | 0 | NA NA | | 20 | | | |
| | N-Nitrosodiphenylamine | | 5 | | | NA 25 | 28 | | | |
| (continued) | Pentachlorophenol | 5 | 5 | 5 | 110 | 35 | 360 | | | |
| - | Phenol | 5 | 5 | 0 | NA 1222 | NA 10 | 420 | | | |
| | 2-Methylnaphthalene | 5 | 5 | 5 | 1200 | 19 | 670 | | 1 | |
| - | 1,1,1,2-Tetrachloroethane | 8 | 9 | 0 | NA | NA | | | | |
| | 1,1,1-Trichloroethane | 8 | 9 | 0 | NA | NA | 856 | | | |
| | 1,1,2 - Trichlorotrifluoroethane | 8 | 9 | 0 | NA | NA | | | | |
| | 1,1,2,2-Tetrachloroethane | 8 | 9 | 0 | NA | NA | 202 | | | |
| | 1,1,2-Trichloroethane | 8 | 9 | 0 | NA | NA | 570 | | | |
| | 1,1-Dichloroethane | 8 | 9 | 0 | NA | NA | 0.575 | | | |
| | 1,1-Dichloroethene | 8 | 9 | 0 | NA | NA | 2780 | | | |
| | 1,1-Dichloropropene | 3 | 4 | 0 | NA | NA | | | | |
| | 1,2,3-Trichlorobenzene | 8 | 9 | 0 | NA | NA | 858 | | | |
| | 1,2,3-Trichloropropane | 8 | 9 | 0 | NA | NA | | | | |
| | 1,2,4-Trichlorobenzene | 8 | 9 | 0 | NA | NA | 31 | | | |
| | 1,2,4-Trimethylbenzene | 8 | 9 | 4 | 980 | 2.4 | | | | |
| F | 1,2-Dibromo-3-chloropropane | 8 | 9 | 0 | NA | NA | | | | |
| | 1,2-Dibromoethane (EDB) | 8 | 9 | 0 | NA | NA | | | | |
| | 1,2-Dichlorobenzene | 8 | 9 | 0 | NA | NA | 35 | | | |
| | 1,2-Dichloroethane (EDC) | 8 | 9 | 0 | NA | NA | 260 | | | |
| - | 1,2-Dichloropropane | 8 | 9 | 0 | NA | NA | 333 | | | |
| - | 1,3,5-Trimethylbenzene | 1 | 1 | 1 | 21 | 21 | 333 | | | |
| | 1,3-Dichlorobenzene | 5 | 5 | 0 | NA NA | NA NA | 842 | | | |
| | 1,3-Dichloropropane | 3 | 4 | 0 | NA NA | NA | 042 | | | |
| | 1,4-Dichloro-2-Butene | 3 | 4 | 0 | NA NA | NA NA | | | | |
| vocs | 1,4-Dichlorobenzene | 2 | 2 | 2 | 23 | 22 | 110 | | | |
| Vocs | 2,2-Dichloropropane | 3 | 4 | 0 | NA | NA | 110 | | | |
| - | 2-Butanone | 8 | 9 | 0 | NA NA | NA NA | 42.4 | | | |
| - | | 3 | 4 | 0 | | | 42.4 | | | |
| - | 2-Chloroethyl Vinyl Ether | | | | NA NA | NA | | | | |
| - | 2-Chlorotoluene | 3 | 4 | 0 | NA | NA | | | | |
| - | 2-Hexanone | 8 | 9 | 0 | NA | NA | 58.2 | | | |
| | 4-Chlorotoluene | 3 | 4 | 0 | NA | NA | | | | |
| - | 4-Methyl-2-pentanone | 8 | 9 | 0 | NA | NA | 25.1 | | | |
| | Acrolein | 3 | 4 | 0 | NA | NA | 0.00152 | | | |
| | Acrylonitrile | 3 | 4 | 0 | NA | NA | 1.2 | | | |
| | Benzene | 8 | 9 | 3 | 8.1 | 1.5 | 137 | | | |
| L | Bromobenzene | 3 | 4 | 0 | NA | NA | | | | |
| L | Bromochloromethane | 8 | 9 | 0 | NA | NA | | | | |
| Ĺ | Bromodichloromethane | 8 | 9 | 0 | NA | NA | | | | |
| | Bromoethane | 3 | 4 | 0 | NA | NA | | | | |
| Γ | Bromoform | 8 | 9 | 0 | NA | NA | 1310 | | | |
| | Bromomethane | 8 | 9 | 0 | NA | NA | 1.37 | | | |
| | Carbon disulfide | 8 | 9 | 1 | 4.3 | 4.3 | 0.851 | | 1* | |
| | Carbon tetrachloride | 8 | 9 | 0 | NA | NA | 7240 | | | |
| ľ | Chlorobenzene | 8 | 9 | 0 | NA | NA | 162 | | | |
| ļ | Chloroethane | 8 | 9 | 0 | NA | NA | | | | |
| | Chloroform | 8 | 9 | 0 | NA | NA | 121 | | | |
| | Chloromethane | 8 | 9 | 0 | NA | NA | | | | |
| F | cis-1,2-Dichloroethene (DCE) | 8 | 9 | 0 | NA | NA | | | 1 | |

Table 3-12 - Summary of Sediment Data

Bremerton Gas Works Superfund Site Bremerton, Washington

| Chemical Group | Chemical Constituent | Number of Locations | Number of Samples | Number of Detections | Maximum Detected Concentration (ug/kg) | Minimum Detected Concentration (ug/kg) | Sediment PRG (ug/kg) | Puget Sound Background Sediment Concentration ¹ (ug/kg) | Number of Detected Concentrations Exceeding the PRG | Number of Detected Concentrations Exceeding Puget Sound Background Metals Concentration |
|----------------|--------------------------------|------------------------|----------------------|-------------------------|---|---|----------------------------|--|---|---|
| | cis-1,3-Dichloropropene | 8 | 9 | 0 | NA | NA | | | | |
| | Cyclohexane | 5 | 5 | 0 | NA | NA | | | | |
| | Dibromochloromethane | 8 | 9 | 0 | NA | NA | | | | |
| | Dibromomethane | 3 | 4 | 0 | NA | NA | | | | |
| | Dichlorodifluoromethane | 5 | 5 | 0 | NA | NA | | | | |
| | Ethylbenzene | 8 | 9 | 2 | 24 | 2.3 | 305 | | | |
| <u> </u> | Hexachlorobutadiene | 8 | 9 | 0 | NA | NA | 11 | | | |
| | Hexachloroethane | 3 | 3 | 0 | NA | NA | 804 | | | |
| | Isopropylbenzene | 8 | 9 | 2 | 9 | 0.48 | 86 | | | |
| | Methyl acetate | 5 | 5 | 0 | NA | NA | | | | |
| | Methyl tert-butyl ether (MTBE) | 5 | 5 | 0 | NA | NA | | | | |
| | Methylcyclohexane | 5 | 5 | 1 | 0.65 | 0.65 | | | | |
| | Methylene chloride | 8 | 9 | 1 | 1.8 | 1.8 | 159 | | | |
| vocs | Methyliodide | 3 | 4 | 0 | NA | NA | | | | |
| | n-Butylbenzene | 3 | 4 | 1 | 84 | 84 | | | | |
| (continued) | n-Propylbenzene | 3 | 4 | 1 | 8.3 | 8.3 | | | | |
| | p-Isopropyltoluene | 3 | 4 | 0 | NA | NA | | | | |
| | sec-Butylbenzene | 3 | 4 | 0 | NA | NA | | | | |
| [| Styrene | 8 | 9 | 0 | NA | NA | 7070 | | | |
| | tert-Butylbenzene | 3 | 4 | 0 | NA | NA | | | | |
| | Tetrachloroethene (PCE) | 8 | 9 | 0 | NA | NA | 190 | | | |
| | Toluene | 8 | 9 | 2 | 1.5 | 0.51 | 1090 | | | |
| [| trans-1,2-Dichloroethene | 8 | 9 | 0 | NA | NA | 1050 | | | |
| | trans-1,3-Dichloropropene | 8 | 9 | 0 | NA | NA | | | | |
| Ī | Trichloroethene (TCE) | 8 | 9 | 0 | NA | NA | 8950 | | | |
| | Trichlorofluoromethane | 8 | 9 | 0 | NA | NA | | | | |
| | Vinyl acetate | 3 | 4 | 0 | NA | NA | 13 | | | |
| | Vinyl chloride | 8 | 9 | 0 | NA | NA | 202 | | | |
| | m,p-Xylenes | 8 | 9 | 2 | 2.9 | 1.7 | | | | |
| [| o-Xylene | 8 | 9 | 2 | 5.7 | 3.9 | | | | |
| [| Naphthalene | 46 | 61 | 61 | 1700000 | 5.4 | 2100 | | 23 | |

Notes:

cPAHs = carcinogenic polycyclic aromatic hydrocarbons

HPAH = high molecular weight PAH

LPAH = low molecular weight PAH

NA = Not applicable, as there are no detections.

PAHs = polycyclic aromatic hydrocarbons

 ${\sf PCBs} = {\sf polychlorinated\ biphenyls}$

PRG = Preliminary Remediation Goal

SVOCs = semivolatile organic compounds

TPH = total petroleum hydrocarbons VOCs = volatile organic compounds

ug/kg = micrograms per kilogram

^{*}Carbon disulfide is a common laboratory chemical. Based on the review of existing analytical data quality, these detections are considered to be the result of laboratory cross-contamination. The results are not considered representative of site conditions.

¹ Background concentrations based on Puget Sound (when available) or Washington State background (Ecology 1994) and the Sediment Cleanup Users Manual II Table 10-1 (Ecology 2015).

Table 4-1 - Summary of Common Ecological Receptors Potentially Present in Vicinity of the Site Bremerton Gas Works Superfund Site

| Common Regional Species | Potentially Use of Site (Yes/Unlikely) | Notes | Reference |
|--|--|---|--|
| quatic Invertebrates | | <u>I</u> | |
| Benthic Invertebrates | | | |
| Amphipods | Yes | | KiTSA 2012 |
| Barnacles | Yes | | GeoEngineers 2011; |
| Benthic Infaunal Community | Yes | | KiTSA 2012 WAC 173-204 |
| Brittle stars | Yes | | GeoEngineers 2011 |
| | | | Anchor QEA 2012; KiT |
| Clams (multiple species) | Yes | | 2012 |
| Mussels (blue and bay) | Yes | | GeoEngineers 2011; KiTSA 2012 |
| Oysters | Yes | | KiTSA 2012 |
| Polycheate worms | Yes | | GeoEngineers 2011; KiTSA 2012 |
| Scallops | Yes | | KiTSA 2012 |
| Sand dollar larvae | Yes | | 1 |
| Sea cucumber | Yes | | |
| Benthivorous Shellfish | | | |
| Octopus | Yes | | KiTSA 2012 |
| <u>Crabs</u> | Yes | | |
| arine-Dependent Birds | | | |
| Piscivorous Raptor Bald eagle | Yes | | KiTSA 2012 |
| <u>Osprey</u> | Yes | Monitored species (state). Nest in Sinclair Inlet. | N13A 2012 |
| Shore Birds | , , , , | | l |
| Belted kingfisher | Yes | | Buchanan 2006 |
| Ducks | Yes | | |
| Glaucous-winged gull | Yes | | KiTSA 2012 |
| Great blue heron | Yes | Monitored species (state). There is a heron rookery along southern Sinclair Inlet (KiTSA 2012). | KiTSA 2012 |
| Marbled murrelet | Unlikely | Threatened (NMFS). Listed marbled murrelet are unlikely to be frequently present in Dyes Inlet (Anchor QEA 2012). | GeoEngineers 2011; Anchor QEA 2012; Ki ⁻ 2012 |
| <u>Sandpiper</u> | Yes | | Buchanan 2006 |
| merganser, northern pintail, parasitic jaeger, plover, red-necked phalarope, rhinoceros auklet, sanderling, <u>sand piper</u> , scaup, scoter, surfbird, tern, turnstone, and American wigeon) | | | 2006 |
| sh | | | |
| Benthivorous Fish Eelpout | Yes | | NOAA 2000 |
| Flatfish (English sole, butter sole, dover sole, sand sole, rock | | | |
| sole, CO sole, and starry flounder) Other bottomfish (skate, sablefish, greenlings, wolf-eel, Pacific | Yes Yes | | KiTSA 2012 NOAA 2000 |
| sanddab, and plainfin midshipman) | | | 11024 2000 |
| Perch (pile and striped) Plainfin midshipman | Yes Yes | | - |
| Poacher | Yes | | - |
| Prickleback | Yes | | 1 |
| Rock sole | Yes | | KiTSA 2012 |
| Spotted ratfish | Yes | | |
| Omnivorous Fish | | | |
| Baby goby | Yes | | NOAA 2000 |
| Chum salmon | Yes | Chum are anadromous and may utilize the site for only a portion of the year. | KiTSA 2012 |
| Coho salmon | Yes | Coho salmon are anadromous and may utilize the site for only a portion of the year. | 2012 Anchor QEA 2012; KI |
| Cutthroat trout | Yes | Cutthroat trout are anadromous and may utilize the site for only a portion of the year. | GeoEngineers 2011; Anchor QEA 2012 |
| Green sturgeon | Unlikely | Threatened (Southern DPS; NMFS). Unlikely to be found in Dyes Inlet (Anchor QEA 2012). | Anchor QEA 2012 |
| Gunnel | Yes | | NOAA 2000 |
| Herring | Yes | Dyes Inlet supports a small herring stock (Anchor QEA 2012). | Anchor QEA 2012 |
| Pink salmon | Yes | Pink salmon are anadromous and may utilize the site for only a portion of the year. | KiTSA 2012 |
| Sockeye salmon | Yes | | |
| Steelhead trout | Yes | Threatened (Puget Sound DPS4; NMFS). Listed Steelhead are anadromous and may utilize the site for only a portion of the year. | GeoEngineers 2011; Anchor QEA 2012 |
| Sculpin (cabezon, Pacific stagehorn, and roughback) | Yes | | NOAA 2000 |
| Sand lance | Yes | May serve as prey to salmonids. | GeoEngineers 2011; Anchor QEA 2012 GeoEngineers 2011; |
| | | | |

Table 4-1 - Summary of Common Ecological Receptors Potentially Present in Vicinity of the Site Bremerton Gas Works Superfund Site

| Common Regional Species | Site (Yes/Unlikely) | Notes | Reference |
|---|--|--|--|
| Piscivorous Fish | | | |
| Bocaccio rockfish | Unlikely | Endangered (Puget Sound/Georgia Basin DPS; NMFS). Rarely observed in Puget Sound (Anchor QEA 2012). | GeoEngineers 2011; Anchor QEA 2012; KiTS 2012 |
| Bull trout | Unlikely | Threatened (Coastal-Puget Sound DPS4; USFWS). Listed bull trout are anadromous. No bull trout stocks have been identified in any of the streams draining into the larger Sinclair Inlet basin, and no designated critical habitat is present within Kitsap County (Anchor QEA 2012). | |
| Canary rockfish | Unlikely | Threatened (Puget Sound/Georgia Basin DPS; NMFS). Unlikely to be present at the site due to unsuitable habitat (Anchor QEA 2012). | |
| Chinook salmon | Yes | Threatened (Puget Sound ESU3; NMFS). Adult Chinook are anadromous and may utilize the site for only a portion of the year. | GeoEngineers 2011; Anchor QEA 2012; KiTS 2012 |
| Ling cod | Yes | | NOAA 2000 |
| Non-listed rockfish (brown, copper, greeenstriped, yellowtail, | Yes | | |
| quillback, black, and yelloweye) | | | |
| Spiny dogfish Yellow rockfish | Yes Unlikely | Threatened (Puget Sound/Georgia Basin DPS; NMFS). Unlikely to be present at the site due to unsuitable habitat | GeoEngineers 2011; |
| | | (Anchor QEA 2012). | Anchor QEA 2012 |
| scivorous Mammals and Other Marine Mammals | | | |
| Dall's porpoise | Yes | Puget Sound resident species. | KiTSA 2012 |
| California sea lion | Yes | Seasonal species. | |
| Gray whale | Unlikely | Seasonal species. Has been observed in Sinclair Inlet. | |
| Harbor porpoise | Yes | Species of concern (state). Puget Sound resident species. Species of concern (state). Puget Sound resident species. | |
| <u>Harbor seal</u> | Yes | Harbor seals are known to be present in Dyes Inlet (Anchor QEA 2012). | |
| Humpback whale | Unlikely | Endangered (NMFS). Humpback whales are infrequently observed in Puget Sound (GeoEngineers 2011). Unlikely to be found in Dyes Inlet (Anchor QEA 2012). | GeoEngineers 2011; Anchor QEA 2012 |
| Killer whale | Unlikely | Endangered (Southern Resident DPS4; NMFS). Listed Orca whales are only present in Puget Sound for a portion of the year (fall/winter). They have been infrequently observed in Dyes Inlet (Anchor QEA 2012). | Anchor QEA 2012; KiT\$ 2012 |
| Minke whale | Unlikely | | KiTSA 2012 |
| Northern sea lion | Yes | Seasonal species. | |
| River otter | Yes | Puget Sound resident species. Risk to species will be addressed by assessment of piscivorous mammal receptor. | |
| Stellar sea lion | Unlikely | Unlikely to be found in Dyes Inlet (Anchor QEA 2012). Risk to species will be addressed by assessment of piscivorous mammal receptor. | GeoEngineers 2011; Anchor QEA 2012; KiT\$ 2012 |
| acrophytes | | | |
| Almos and Irela | Yes | Aquatic vegetation in Dyes Inlet is patchy (Anchor QEA 2012). Dyes Inlet and Sinclair Inlet do not support any floating kelp (Anchor QEA 2012). Non-floating kelp species are present in | |
| Algae and kelp | | just 18% of the shoreline throughout the entire basin (PSP 2005; Anchor QEA 2012). May serve as habitat and food for marine species (KiTSA 2012). | 2012 |
| Algae and kelp Popweed | Yes | 2005; Anchor QEA 2012). May serve as habitat and food for | 2012 KiTSA 2012 |
| | Yes Yes | 2005; Anchor QEA 2012). May serve as habitat and food for | |
| Popweed | | 2005; Anchor QEA 2012). May serve as habitat and food for | |
| Popweed Sea lettuce | Yes | 2005; Anchor QEA 2012). May serve as habitat and food for marine species (KiTSA 2012). Within Dyes Inlet and Chico Bay there are scattered patches of eelgrass in intertidal areas with muddy to sandy substrates (WDNR 2001; Anchor QEA 2012). Suitable eelgrass habitat | KiTSA 2012 Anchor QEA 2012; KiT: |
| Popweed Sea lettuce Eelgrass | Yes | 2005; Anchor QEA 2012). May serve as habitat and food for marine species (KiTSA 2012). Within Dyes Inlet and Chico Bay there are scattered patches of eelgrass in intertidal areas with muddy to sandy substrates (WDNR 2001; Anchor QEA 2012). Suitable eelgrass habitat | KiTSA 2012 Anchor QEA 2012; KiTs |
| Popweed Sea lettuce Eelgrass | Yes | 2005; Anchor QEA 2012). May serve as habitat and food for marine species (KiTSA 2012). Within Dyes Inlet and Chico Bay there are scattered patches of eelgrass in intertidal areas with muddy to sandy substrates (WDNR 2001; Anchor QEA 2012). Suitable eelgrass habitat | KiTSA 2012 Anchor QEA 2012; KiTs |
| Popweed Sea lettuce Eelgrass rrestrial Species Avian Predator Black-capped Chickadee Crow | Yes Unlikely Yes Yes | 2005; Anchor QEA 2012). May serve as habitat and food for marine species (KiTSA 2012). Within Dyes Inlet and Chico Bay there are scattered patches of eelgrass in intertidal areas with muddy to sandy substrates (WDNR 2001; Anchor QEA 2012). Suitable eelgrass habitat | KiTSA 2012 Anchor QEA 2012; KiT: 2012 |
| Popweed Sea lettuce Eelgrass rrestrial Species Avian Predator Black-capped Chickadee Crow Evening grosbeak | Yes Unlikely Yes Yes Yes Yes | 2005; Anchor QEA 2012). May serve as habitat and food for marine species (KiTSA 2012). Within Dyes Inlet and Chico Bay there are scattered patches of eelgrass in intertidal areas with muddy to sandy substrates (WDNR 2001; Anchor QEA 2012). Suitable eelgrass habitat | KiTSA 2012 Anchor QEA 2012; KiT: 2012 |
| Popweed Sea lettuce Eelgrass Prestrial Species Avian Predator Black-capped Chickadee Crow Evening grosbeak Flicker | Yes Unlikely Yes Yes Yes Yes Yes Yes | 2005; Anchor QEA 2012). May serve as habitat and food for marine species (KiTSA 2012). Within Dyes Inlet and Chico Bay there are scattered patches of eelgrass in intertidal areas with muddy to sandy substrates (WDNR 2001; Anchor QEA 2012). Suitable eelgrass habitat | KiTSA 2012 Anchor QEA 2012; KiT 2012 |
| Popweed Sea lettuce Eelgrass Prestrial Species Avian Predator Black-capped Chickadee Crow Evening grosbeak Flicker Golden-crowned kinglet | Yes Unlikely Yes Yes Yes Yes Yes Yes Yes | 2005; Anchor QEA 2012). May serve as habitat and food for marine species (KiTSA 2012). Within Dyes Inlet and Chico Bay there are scattered patches of eelgrass in intertidal areas with muddy to sandy substrates (WDNR 2001; Anchor QEA 2012). Suitable eelgrass habitat is not present at the Site. | KiTSA 2012 Anchor QEA 2012; KiT 2012 |
| Popweed Sea lettuce Eelgrass Prestrial Species Avian Predator Black-capped Chickadee Crow Evening grosbeak Flicker Golden-crowned kinglet Purple martin | Yes Unlikely Yes Yes Yes Yes Yes Yes Yes Yes | 2005; Anchor QEA 2012). May serve as habitat and food for marine species (KiTSA 2012). Within Dyes Inlet and Chico Bay there are scattered patches of eelgrass in intertidal areas with muddy to sandy substrates (WDNR 2001; Anchor QEA 2012). Suitable eelgrass habitat | KiTSA 2012 Anchor QEA 2012; KiT 2012 |
| Popweed Sea lettuce Eelgrass Frrestrial Species Avian Predator Black-capped Chickadee Crow Evening grosbeak Flicker Golden-crowned kinglet Purple martin Ring-necked pheasant | Yes Unlikely Yes Yes Yes Yes Yes Yes Yes Yes Yes Ye | 2005; Anchor QEA 2012). May serve as habitat and food for marine species (KiTSA 2012). Within Dyes Inlet and Chico Bay there are scattered patches of eelgrass in intertidal areas with muddy to sandy substrates (WDNR 2001; Anchor QEA 2012). Suitable eelgrass habitat is not present at the Site. | KiTSA 2012 Anchor QEA 2012; KiT 2012 |
| Popweed Sea lettuce Eelgrass Prestrial Species Avian Predator Black-capped Chickadee Crow Evening grosbeak Flicker Golden-crowned kinglet Purple martin Ring-necked pheasant Robin | Yes Unlikely Yes Yes Yes Yes Yes Yes Yes Yes Yes Ye | 2005; Anchor QEA 2012). May serve as habitat and food for marine species (KiTSA 2012). Within Dyes Inlet and Chico Bay there are scattered patches of eelgrass in intertidal areas with muddy to sandy substrates (WDNR 2001; Anchor QEA 2012). Suitable eelgrass habitat is not present at the Site. | KiTSA 2012 Anchor QEA 2012; KiT 2012 |
| Popweed Sea lettuce Eelgrass Prestrial Species Avian Predator Black-capped Chickadee Crow Evening grosbeak Flicker Golden-crowned kinglet Purple martin Ring-necked pheasant Robin Starling | Yes Unlikely Yes Yes Yes Yes Yes Yes Yes Yes Yes Ye | 2005; Anchor QEA 2012). May serve as habitat and food for marine species (KiTSA 2012). Within Dyes Inlet and Chico Bay there are scattered patches of eelgrass in intertidal areas with muddy to sandy substrates (WDNR 2001; Anchor QEA 2012). Suitable eelgrass habitat is not present at the Site. | KiTSA 2012 Anchor QEA 2012; KiT 2012 |
| Popweed Sea lettuce Eelgrass Prestrial Species Avian Predator Black-capped Chickadee Crow Evening grosbeak Flicker Golden-crowned kinglet Purple martin Ring-necked pheasant Robin Starling Steller's jay | Yes Unlikely Yes Yes Yes Yes Yes Yes Yes Yes Yes Ye | 2005; Anchor QEA 2012). May serve as habitat and food for marine species (KiTSA 2012). Within Dyes Inlet and Chico Bay there are scattered patches of eelgrass in intertidal areas with muddy to sandy substrates (WDNR 2001; Anchor QEA 2012). Suitable eelgrass habitat is not present at the Site. | KiTSA 2012 Anchor QEA 2012; KiT 2012 |
| Popweed Sea lettuce Eelgrass Prestrial Species Avian Predator Black-capped Chickadee Crow Evening grosbeak Flicker Golden-crowned kinglet Purple martin Ring-necked pheasant Robin Starling Steller's jay Carnivorous Mammals | Yes Unlikely Yes Yes Yes Yes Yes Yes Yes Yes Yes Ye | 2005; Anchor QEA 2012). May serve as habitat and food for marine species (KiTSA 2012). Within Dyes Inlet and Chico Bay there are scattered patches of eelgrass in intertidal areas with muddy to sandy substrates (WDNR 2001; Anchor QEA 2012). Suitable eelgrass habitat is not present at the Site. | KiTSA 2012 Anchor QEA 2012; KiT 2012 KiTSA 2012 |
| Popweed Sea lettuce Eelgrass Prestrial Species Avian Predator Black-capped Chickadee Crow Evening grosbeak Flicker Golden-crowned kinglet Purple martin Ring-necked pheasant Robin Starling Steller's jay | Yes Unlikely Yes Yes Yes Yes Yes Yes Yes Yes Yes Ye | 2005; Anchor QEA 2012). May serve as habitat and food for marine species (KiTSA 2012). Within Dyes Inlet and Chico Bay there are scattered patches of eelgrass in intertidal areas with muddy to sandy substrates (WDNR 2001; Anchor QEA 2012). Suitable eelgrass habitat is not present at the Site. | KiTSA 2012 Anchor QEA 2012; KiT 2012 |

Table 4-1 - Summary of Common Ecological Receptors Potentially Present in Vicinity of the Site

Bremerton Gas Works Superfund Site Bremerton, Washington

| Common Regional Species | Potentially use of Site (Yes/Unlikely) | Notes | Reference |
|---------------------------|--|---|---|
| Herbivorous Mammals | (1.00/0 | 110100 | - Kalaranaa |
| Deer | Unlikely | Unlikely to be present at the site due to unsuitable habitat. | KiTSA 2012 |
| Rabbits | Yes | | 7 |
| Squirrels | Yes | | |
| Vole | Yes | | |
| Insectivorous Mammal | | | |
| Shrews | Yes | | |
| Omnivorous Mammals | | T | <u></u> |
| Black bear | Unlikely | Unlikely to be present at the site due to unsuitable habitat. | KiTSA 2012 |
| Mice | Yes | | |
| Moles | Yes | | |
| Raccoon | Yes | | |
| Other Miscellaneous Fauna | | The base of the second | <u> </u> |
| Garter snakes | Yes | Habitat at the site includes the upland embankment and unpaved upland site areas. | KiTSA 2012 |
| Newts and frogs | Unlikely | Amphibians are not likely to be present at the site due unsuitable habitat. | |
| Salamanders | Unlikely | Amphibians are not likely to be present at the site due unsuitable habitat. | |
| Turtles | Unlikely | Turtles are not likely to be present at the site due to unsuitable habitat. | |
| Upland Vegetation | | | _ |
| Big leaf maple | Yes | Native vegetation. Limited in developed site areas. | KiTSA 2012 |
| Douglas fir | Yes | | GeoEngineers 2011; KiTSA 2012 |
| Kinnikinnick | Yes | | KiTSA 2012 |
| Oregon grape | Yes | - | |
| Pacific madrone | Yes | 4 | |
| Pacific rhododendron | Yes Yes | - | Coo Engineero 2011 |
| Pacific gumweed | | 1 | GeoEngineers 2011 GeoEngineers 2011; |
| Red alder | Yes | | Anchor QEA 2012 |
| Salal | Yes | | KiTSA 2012 |
| Sword fern | Yes | | |
| Vine maple | Yes | | |
| Western hemlock | Yes | | |
| Western red cedar | Yes | | |
| Japanese knotweed | Yes | Non-native species. | Anchor QEA 2012 |
| Himalayan blackberry | Yes | | GeoEngineers 2011; Anchor QEA 2012; KiTS 2012 |
| Magnolia | Yes | 1 | GeoEngineers 2011 |
| Pampas grass | Yes |] | |
| Scotch broom | Yes | | GeoEngineers 2011; KiTSA 2012 |
| Spear saltbrush | Yes | | GeoEngineers 2011 |
| Thistle | Yes | | KiTSA 2012 |

Notes:

<u>Underlined = Representative species included as part of ecological A31CSM figures.</u>

 $\label{eq:continuous} \mbox{Anchor QEA, 2012. } \mbox{\it Biological Evaluation} \; . \; \; \mbox{Chico Creek Estuary Restoration Project.} \; \; \mbox{\it January 2012.} \; \mbox{\it Chico Creek Estuary Restoration Project.} \; \mbox{\it January 2012.} \; \mbox{\it Chico Creek Estuary Restoration Project.} \; \mbox{\it January 2012.} \; \mbox{\it Chico Creek Estuary Restoration Project.} \; \mbox{\it Ch$

Buchanan, J.B., 2006. *Nearshore Birds in Puget Sound*. Puget Sound Nearshore Partnership. Report number 2006-05. Published by Seattle District, U.S. Army Corps of Engineers, Seattle, Washington.

DPS = distinct population segment

ESU = evolutionary significant unit

GeoEngineers, 2011. Biological Assessment. Bay Street Pedestrian Enhancement/Mosquito Fleet Trail Project. LSTPE-0166 (008). Port Orchard, Washington. Prepared for City of Port Orchard. August 26, 2011.

KiTSA (Kitsap Trees and Shoreline Association), 2012. Sinclair Inlet Development Concept Plan. Sponsored by KiTSA.

NMFS = National Marine Fisheries Service

NOAA (National Oceanic and Atmospheric Administration), 2000. Gustafson R.G., W.H. Lenarz, B.B. McCain, C.C. Schmitt, W.S. Grant, T.L. Builder, and R.D. Methot. 2000. Status review of Pacific Hake, Pacific Cod, and Walleye Pollock from Puget Sound, Washington. U.S. Dept. Commer., NOAA Tech. Memo. NMFS-NWFSC- 44, 275 p. PSP (Puget Sound Partnership), 2005. Regional Nearshore and Marine Aspects of Salmon Recovery. June 2005.

USFWS = U.S. Fish and Wildlife Service

WAC = Washington Administrative Code

WDNR (Washington Department of Natural Resources), 2001. WDNR and Puget Sound Assessment and Monitoring Program Shore Zone data set. Nearshore Habitat Program. Threatened and endangered species will be re-evaluated at the time of the risk assessment.

Table 4-2 - Nationwide MGP Site Summary Bremeton Gas Works Superfund Site Bremerton, Washington

| MGP Site Name & Location | Reference | Geologic Conditions | Groundwater / Surfacewater | Chemicals of Concern | Remedial Actions | Cleanup Status |
|--|---|---|--|-------------------------|--|------------------------------|
| Cold Spring MGP Site Cold Spring, NY | Record of Decition (2010) http://www.dec.ny.gov/docs/r emediation_hudson_pdf/e34 0026arod.pdf *Subsurface soils consist of 11-13 feet of debris containing fill underlain by a 15 foot thick layer of c which overfies bedrock. *Contamination confined to the fill material. | | *Groundwater flows to the west, towards the Hudson River which is adjacent to the site. No contamination was observed in river sediments. *No Hudson BTEX PAHs | | Excavation and off-site treatment/disposal. | Scheduled to begin late 2014 |
| Saranac Street MGP Site Plattsburgh, NY | http://www.dec.ny.gov/docs/r emediation_hudson_pdf/rod 51000701.pdf | Subsurface soils consist of up to 21 feet of debris containing fill undertain by up to 15 foot thick layer of sandy alluvium. Beneith the alluvium lies a layer of dense glacial till, which overlies limestone bedrock. Contamination present down to and into fractured bedrock. | derlain by up to 15 foot thick layer of Beneith the alluvium lies a layer of , which overlies limestone bedrock. •Coal tar discharged into the river along the | | In situ stabilization; Soil and sediment excavation with off-site treatment/disposal; Bedrock tar collection wells. | Remedial Action complete |
| Waterville MFG Plant Waterville, NY | http://www.dec.ny.gov/docs/r emediation hudson pdf/6330 41_1.pdf | Subsurface soils consist of one foot of topsoil over a fill unit up to 12 feet thick consisting of a substantial amount of ash as well as brown sand and gravel, coal fragments and bricks. Below the fill is a unit of glacial outwash sand and silt ranging in thickness from 1 to 10 feet. A dense kame moraine silt and gravel deposit of depths from 4 to 12 feet was found below the outwash unit. *Contamination present up to 14 feet below grade. | A western flowing tributary to Big Creek forms the southern edge of the property, approximately 150 feet south of the site. The depth to groundwater ranges from approximately 4 to 12 feet below grade. Groundwater flow through the site is to the south-southwest and discharges into the Big Creek tributary. | BTEX PAHs | Excavation and Disposal; Institutional Controls; Soil Cap. | No Further Action required |
| Cortland Homer Former MGP Site Homer, NY | mttp://www.dec.ny.gov/docs/r | Subsurface soils consist of a fill layer ranging from 6 inches to 10 feet and is underlain by outwash sand that varies in thickness from 20 to 40 feet. A confining silf/clay layer was observed benieth the outwash sand. Contamination present up to 37 feet below grade. | The West Branch of the Tioughnioga River is located 150 feet east of the site parcels. Depth to groundwater at the site is approximately 5 feet below grade. Groundwater flow is in a east to east-southeast direction. Groundwater discharges into the river. River sediments have been impacted by contaminants. | BTEX PAHs Cyanide | Excavation and disposal of source area soils; In situ stabilization of downgradient contaminated soils; NAPL collection trench; Sediment removal. | Remedial Design complete |

Table 4-2 - Nationwide MGP Site Summary Bremeton Gas Works Superfund Site Bremerton, Washington

| MGP Site Name & Location | Reference | Geologic Conditions | Groundwater / Surfacewater | Chemicals of Concern | Remedial Actions | Cleanup Status |
|--|---|---|--|----------------------|---|--|
| Tacoma Tar Pits Tacoma, WA | http://yosemite.epa.gov/R10/ CLEANUP.NSF/sites/TacomaTa rpis/SFILE/TTP-SYr-Review_ Sept03.pdf | *Subsurface soils consist of several feet of fill underlain by a layered sequence of silts and sands. | The Puyallup River is just norheast of the site. Groundwater occurs several feet below ground surface at the Tacoma Tar Pits site. The groundwater levels at the site vary in response to the tidal action in Commencement Bay and adjacent waterways. Groundwater flow directions vary depending on location, season, and tide stage. In general however, groundwater typically flows east (northwest and central potions of the site) and south (southeast portion of the site). | BTEX PAHs | Excavation and stabilization; Stabilized material placed in an engineered waste pile on site; Soil cap; Groundwater pump and treat. | Ongoing O&M for cover and groundwater treatment system |
| Oakland MGP Oakland, CA | | Subsurface soils consisting of up to 5 feet of gravel/sand fill underlain by a sandy layer that extends up to 15 feet below grade with interbeded layers of silt and clay. The sandy layer is underlain by a fine-grained layer of clay and silt up to 20 feet below grade. Contamination present up to 21 feet below grade. | TPH Groundwater is 2 to 7.5 feet bgs and flows towards the akland Inner Harbor, which is approximately 1000 feet way. PAHs Cyanide | | Soil cap. | Ongoing O&M |
| Glens Falls - Mohican Street MGP Glens Falls, NY | http://www.dec.ny.gov/docs/r emediation hudson pdf/5570 16roda2.pdf | Subsurface soil cosists of fill underlain by glacial fluvial deposits of sand, slit, silty sand, sandy slit. A layer of silty day overlies bedrock, which is encountered between 9-29 feet below grade. Contamination present up to 19 feet below grade. | The site is bounded to the south by the Glens Falls feeder canal. Groundwater is 2-14 feet below grad and flows towards the Glens Falls canal and Hudson River. Canal sediments are impacted. | BTEX PAHs | Excavation of source material; Oxygen delivery system; Soil cover; Institutional controls; Dredging and disposal. | Remedial Action approved |
| Gastown MGP Site Tonawanda, NY | http://www.dec.ny.gov/docs/r emediation hudson.pdf/rod9 15171text.pdf http://www.dec.ny.gov/chemi cal/\$8387.html | Subsurface soils consist of up to 22 feet of debris containing fill undertaini by layers of sand and silt for an additional 24 feet below grade. Contamination present down into the sand/silt layers. | The site is bounded to the north-northwest by Tonawanda Creek. Groundwater is approximately 6 feet below grade and flows to the north into Tonawanda Creek. Creek sediments have been impacted. | BTEX PAHs | Excavation and disposal; In situ stabilization; NAPL collection wells. | Scheduled to begin in 2013 |
| Former Sacramento MGP Sacramento, CA | http://www.pge.com/about/e nvironment/taking- responsibility/mgp/sacramen to.shtml | Subsurface soils consist of up to 15 feet of fill undertain by a layer containing mostly silts and dayey silts to 25 feet below grade. A layer of unconclidated sand extends from approximately 25 feet to 85 feet below grade. Contamination present up to 45 feet below grade. | The site is located adjacent to the Sacramento River. Groundwater is present approximately 18 feet below grade and flow is strongly incluenced by the Sacramento River and flows to the east. | TPH BTEX PAHs | Excavation and disposal; Pump and treat; In stu stabilization. | <i>In situ</i> stabilization implemented late 2012 |
| Former Red Bluff MGP Red Bluff, CA | nvironment/taking- | Subsurface soil consists of up between 3 and 28 feet of debris containing fill material underlain by a sily clay / clayey silt with interbedded sand, grave, and finergrained sediments. Contamination present in the fill material. | The site is bound to the east by the Sacramento River. Groundwater is present between 4 and 39 feet below grade and is heavily influenced by river level. Groundwater flows either east, or west, depending on river stage. | TPH BTEX PAHs | Excavation and disposal of shallow source soils; In situ stabalization of deeper source soils. | Remedial Action approved |

Table 4-2 - Nationwide MGP Site Summary Bremeton Gas Works Superfund Site Bremerton, Washington

| MGP Site Name & Location | Reference | Geologic Conditions | Groundwater / Surfacewater | Chemicals of Concern | Remedial Actions | Cleanup S tatus |
|---|---|--|---|-------------------------|--|---|
| Georgia MGP | m/stuff/contentmgr/files/0/50 | Subsurface soil consists of up to 22 feet of fill underlain by 15 feet of alluvium above weathered bedrock. Contamination present to the bedrock. | The site is bounded to the west by the Chattahoochee River. | BTEX PAHs | In situ stabilization; Excavation and disposal; Groundwater barrier. | Remedial Action complete |
| Nyack MGP Site Nyack, NY | nttp://www.dec.ny.gov/docs/r | Subsurface soil consists of up to 13 feet of fill underlain by native sitly sand and glacial till layers. Sandstone bedrock was encountered approximately 40 feet below grade. Contamination present to the bedrock. | The site is bound to the north by the Hudson River. The bedrock is a productive aquifer with the groundwater flowing upward through the bedrock. Groundwater generally flows toward the Hudson River. River sediments have been impacted. | BTEX PAHs | Excavation and disposal; "In situ stabilization; In situ chemical oxidation; Dredging and disposal. | Upland solidification complete. Sediment removal scheduled to begin in 2013 |
| Manitowoc Former MGP Site Manitowoc, WI | http://www.epa.gov/region05 /cleanup/manitowoc/pdfs/m anitowoc-completion-report- | Subsurface soil consists of 3-10 feet of miscellaneous sand/silt/clay fill material overlying glacial deposits of sind with varying amounts of gravel, silt, and clay. Unconsolidated materials extend to at least 40 feet below grand and bedrock is estimated to be approximately 48 to 50 feet below grade. Contamination present up to 27 feet below grade. | The site is bound to the northwest by the Manitowoc River. Groundwater is present between 5 and 22 feet below grade and flows towards the Manitowoc River. River sediments have been impacted. | BTEX PAHs Cyanide | Shallow excavation and disposal; In situ stabilization; Pump and treat (carbon); In situ stabilization for sediments failed; Dredging. | Pump and Treat O&M Sediment dredging scheduled to begin December 2013 |
| Kinston MGP Site Kinston, NC | http://www.neuselibrary.org/ Kinston%20MGP%20Reme | Subsurface soils consist of gravel fill underlain by a fine to medium grained sand layer with some gravel and clay up to 21 feet below grade. The sandy layer is underlain by a silt/day which extends up to 45 feet below grade, followed by a silty sand extending to 55 feet below grade. Contamination present up to 23 feet below grade. | The Neuse River borders more than 50% of the Site including the north, west, and southwest boundaries. Groundwater flow is to the southwest, towards the Neuse River. River sediments have been impacted. | BTEX PAHs Cyanide | In situ stabalization; Institutional controls. | Remedy selected, awaiting implementation |

BTEX = benzene, tolousne, ethylbenzene, and xylenes

cPAHs = carcinogenic polycyclic aromatic hydrocarbons

MGP = manufactured gas plant

NAPL = non-aqueous phase liquid

O&M = operation and maintenance

PAHs = polycyclic aromatic hydrocarbons TPH = total petroleum hydrocarbons

| | | | | | | | | Potent | ial Sou | rces of | MGP-R | | Contar | ninants | s | Potentia | l Human He | alth and |
|----------------------------------|--|--------------------------------------|------------------------------|---|---------------------------------------|------------------------|--|-----------------|----------------------|--|---|------------------------------|--|---|---|----------------------------------|---|------------------------------------|
| D1 | iminary Contaminants of Potential Concern | | D | on for Incl | | | Feed | dstocks | and | | | | - D | | | Enviro | nmental Co | ncerns |
| Prei | minary Contaminants of Potential Concern | | Reas | on for incit | usion | | | Fuels | | | WIGP | Proces | s Bypro | aucts | (Š | ' | see Note 2 | , |
| Contaminant Group | <u>Contaminant</u> Benzene | × Potential MGP-Related Constituents | × Other Sources (see Note 3) | × Detected in Previous Sampling Efforts | × Detected Above Initial PRGs | Other EPA Contaminants | × Gasoline | Diesel fuel oil | Coal/coke briquettes | Ash, clinker, cinder, slag, soot, bricks | × Spent scrubber media (tarry wood chips) | × Tar (potentially as DNAPL) | × Light oil (potentially as LNAPL) | × Gas Liquor and Emulsion (tar-water mixture) | × Spent purifier media (iron oxide, tarry wood chips) | × Human Health Risk (Carcinogen) | × Other Human Health Risks (non-Carcinogen) | × Toxicity to Ecological Receptors |
| | Toluene | X | X | X | _ ^ | | X | | | | X | X | X | X | X | | X | X |
| | Ethylbenzene | Х | Х | Х | Х | | Х | | | | Х | Х | Х | Х | Х | | Х | Х |
| | Xylenes | Х | Х | X | Х | | Х | | | | Х | Х | Х | Х | Х | | X | Х |
| 1 | 1,2,3-Trichlorobenzene 1,2,4-Trimethylbenzene | х | х | X | | 1 | Х | - | - | \vdash | Х | Х | Х | Х | Х | - | X | х |
| 1 | 1,3,5-Trimethylbenzene | _^ | | X | | | ^ | | | | | <u> </u> | ^ | ^ | | | X | _^_ |
| 1 | 1,4-Dichlorobenzene | | | Х | | | | | | | | | | | | Х | Х | Х |
| 1 | 1,4-Difluorobenzene | | v | X | | - | <u> </u> | _ | - | _ | | | <u> </u> | - | | U. | ,, | |
| | 1,2-Dichloroethane 2-butanone | | Х | X | Х | | | | | | | | | | | Х | X | X |
| | Acetone | | | X | | | | | | | | | | | | | X | X |
| 8 | Carbon disulfide | | | Х | | | | | | | | | | | | | Х | Х |
| Volatile Organic Compounds | Carbon Tetrachloride | | Х | Х | Х | | | | | | | | | | | Х | Х | Х |
| l g | Chlorobenzene-d5 | | . v | X | V | | | - | | - | | | | | | | X | X |
| Ō | Chloroform cis-1,3-Dichloropropene | | X | X | X | | | | | | | | | | | X | X | Х |
| anic | cis-1,2-Dichloroethene | | ^ | X | _ ^ | | | | | | | | | | | ^ | X | |
| Org | Cyclochexane | | | Х | | | | | | | | | | | | | X | |
| i⊨ | Isopropylbenzene | | | Х | | | | | | | | | | | | | Х | Х |
| /ola | Methyl acetate | | | X | | | | | | <u> </u> | | | | | | | Х | |
| _ | Methylcyclohexane | | | X | | | | | | | | | | | | V | V | |
| | Methylene chloride n-Butylbenzene | | | X | | | | | | | | | | | | Х | X | Х |
| | n-Hexane | | | X | Х | | | | | | | | | | | | X | Х |
| | n-Propylbenzene | | | Х | | | | | | | | | | | | | Х | |
| | Pentafluorobenzene | | | Х | | | | | | | | | | | | | | |
| | p-Isopropyltoluene | | | X | | | | | | | | | | | | | v | |
| | sec-Butylbenzene Styrene | | | X | | | | | | | | | | | | | X | Х |
| 1 | Tetrachloroethene | | | X | | | | | 1 | | | | | | | Х | X | X |
| 1 | trans-1,3-Dichloropropene | | Х | X | Х | | | | | | | | | | | X | X | |
| 1 | Trichloroethene | | Х | Х | Х | | | | | | | | | | | Х | Х | Х |
| <u> </u> | Trichlorofluoromethane | ,, | ., | X | \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ | - | | - | ., | L | | \ , | _ | , , | | V. | X | X |
| | Benzo(a) anthracene Benzo(b) fluoranthene | X | X | X | X | - | | - | X | X | X | X | | X | X | X | X | X |
| | Benzo(k)fluoranthene | × | x | × | X | | | | x | × | X | X | | x | X | X | × | X |
| | Benzo(a)pyrene | X | Х | X | X | | L | L | X | X | X | Х | L | Х | X | X | X | X |
| ons | Chrysene | Х | Х | Х | Х | | | | Х | Х | Х | Х | | Х | Х | Х | Х | Х |
| P. P. | Dibenz(a,h)anthracene | X | X | X | X | | <u> </u> | - | X | X | X | X | _ | X | X | X | X | X |
| ĕ | Indeno(1,2,3-cd)pyrene Acenaphthene | X | X | X | X | | - | Х | X | X | X | X | - | X | X | Х | X | X |
| ž | Acenaphthylene | X | X | X | X | | | × | × | × | X | X | | × | X | | X | X |
| atic | Anthracene | X | Х | X | X | | L | Х | Х | Х | Х | Х | L | Х | Х | | X | X |
| 1 m | Benzo(g,h,i)perylene | Х | Х | Х | Х | | | Х | Х | Х | Х | Х | | Х | Х | | Х | Х |
| Polycyclic Aromatic Hydrocarbons | Dibenzofuran | .,, | | X | X | | <u> </u> | L., | L., | L., | ., | L., | - | L., | L., | | X | X |
| رم ارم | Fluoranthene Fluorene | X | X | X | X | - | | X | X | X | X | X | - | X | X | - | X | X |
| ا کا ا | Phenanthrene | X | X | X | X | | | X | x | X | X | X | | X | X | | X | X |
| " | Pyrene | X | X | X | X | | | X | X | X | X | X | | X | X | | X | X |
| | Methylnaphthalene, 1- | Х | Х | Х | Х | | Х | Х | Х | Х | Х | Х | Х | Х | Х | | Х | Х |
| | Methylnaphthalene, 2- | X | Х | X | X | | Х | X | X | X | X | X | X | X | X | | X | X |
| | Naphthalene | Х | Х | Х | Х | <u> </u> | Х | Х | Х | Х | Х | Х | Х | Х | Х | <u> </u> | Х | Х |

Bremeton Gas Works Superfund Site Bremerton, Washington

| Prel | iminary Contaminants of Potential Concern | | Reas | on for Incl | ısion | | | Potent Istocks Fuels | | (see Note 1) MGP Process Byproducts | | • | Enviro | l Human He nmental Co (see Note 2) | ncerns | | | |
|---------------------------------------|---|------------------------------------|----------------------------|---------------------------------------|-----------------------------|------------------------|----------|----------------------------|----------------------|--|---|----------------------------|----------------------------------|---|---|--------------------------------|---|----------------------------------|
| Contaminant Group | Contaminant | Potential MGP-Related Constituents | Other Sources (see Note 3) | Detected in Previous Sampling Efforts | Detected Above Initial PRGs | Other EPA Contaminants | Gasoline | Diesel fuel oil | Coal/coke briquettes | Ash, clinker, cinder, slag, soot, bricks | Spent scrubber media (tarry wood chips) | Tar (potentially as DNAPL) | Light oil (potentially as LNAPL) | Gas Liquor and Emulsion (tar-water mixture) | Spent purifier media (iron oxide, tarry wood chips) | Human Health Risk (Carcinogen) | Other Human Health Risks (non-Carcinogen) | Toxicity to Ecological Receptors |
| Compounds | 1.1'-Biphenyl 1.2,4-Trichlorobenzene 2,4-Dimethylphenol 4-Methylphenol Acetophenone | | | X X X X | | | | | | | | | | | | X | X X X | X X X |
| Other Semi-volatile Organic Compounds | Benzyl butyl phthalate Benzaldehyde Bis(2-ethylhexyl) phthalate Caprolactam Carbazole | X | | X X X | | | | | | | X | Х | | X | X | X | X X X X | X |
| Other Semi-w | creosols Dibenzofuran Di-n-butyl phthalate Isophorone Pentachlorophenol | X | X | X X X | X | | | | | | X | X | | X | X | X X | X X X X | X X X X |
| | Phenol Aluminum Antimony Arsenic Barium | X | X | X X X | X | | | | | | X | X | | X | X | X | X X X X | X X X X |
| S 4 | Beryllium Cadmium Chromium Cobalt Copper | | X X | X X X X | X X X | | | | | | | | | | | | X X X X | X X X X |
| Metals ⁴ | Iron Lead Manganese Mercury Nickel | | X | X X X X | XXX | | | | | | | | | | | | X X X X | X X X X |
| | Selenium Silver Thallium Vanadium | | | X X X | X X | | | | | | | | | | | | X X X | X X X |
| Pestic | Zinc Polychlorinated Biphenyls (PCBs) ⁵ Pesticides ⁶ | | Х | X | X | X X | | | | | | | | | | X X | X X X | X X X |
| Other | s/Furans ⁷ Cyanide, WAD Cyanide, total Sulfide | X X X | | | | X | | | | | | | | | X | X | X X X | X X X |

- 1) Contaminants of Potential Concern (COPCs) associated with MGP sources based on typical composition of MGP-related feedstocks and byproducts (see Section 2.3.1.1)
- 2) Potential Human Health and Environmental Concerns identified based on whether risk-based screening levels or potential ARARs for human health (carinogenic health effects), human health (non-carcinogenic health effects), or ecological health effects were identified during development of initial Preliminary Remediation Goals (PRGs) (see Section 6).
- 3) Other Sources include other historical operations at the site or regional sources of contamination
- 4) Although previously detected at the Site, non-toxic metals (calcium, magnesium, sodium and potassium) are not included herein. Initial PRGs were not developed for these metals because they are essential nutrients that can be tolerated in high doses by living systems.
- 5) PCBs were previously analyzed for and not detected above reporting limits in soil or groundwater at the Site. However, the full standard list of PCB aroclors are COPCs for further evaluation
- 6) The full standard list of pesticides, identified and quantified by EPA Method 8081B, are preliminary COPCs
- 7) The full standard list of dioxins and furans identified and quantified by EPA Method 8290, are preliminary COPCs

This table is not intended to be an exhaustive and complete preliminary list of Site COPCs. The RI/FS will include analysis of samples for the full standard list of analytes under each contaminant group. This list will be evaluated and revised as data is collected and specific contaminants can either be eliminated from the COPC list or are identified as Site COPCs.

Table 5-1 - Summary of Existing Information and Data Gaps - UplandsBremerton Gas Works Superfund Site Bremerton, Washington

| Remdial Investigation/Feasbility Study Information Needs by Topic | Existing Information | Data Gaps | Recommended Data Collection |
|---|--|---|---|
| Physical Characteristics | | | |
| Characteristics of water-bearing zones | Soil stratigraphy and observed/measured groundwater occurrence from previous investigations identifies a waterbearing zone in clean to silty glacial sands at depths of 15 to 41 feet below surface. | Measured/tested physical properties of soil comprising water-bearing zones and aquitards. Hydraulic conductivity of water-bearing zones and aquitards. Vertical extent of the shallow water-bearing zone Presence, location and nature of aquitards. Presence, location and nature of deeper water-bearing zones. | Soil borings to evaluate soil stratigraphy and identify water-bearing zones and aquitards. Soil samples from borings for laboratory measurement of physical parameters that may include grain size, porosity, bulk density, and total/fraction organic carbon. Slug tests at select site wells to measure hydraulic conductivity in each saturated stratigraphic horizon and in different water-bearing zones (if applicable). |
| Groundwater flow direction and gradient | Manual groundwater level measurements collected at eight wells in 2007 were used to evaluate groundwater flow direction and gradient. | Groundwater flow direction and horizontal/vertical gradients. Seasonal variability in water levels and groundwater gradients. Influence of precipitation/surface water infiltration on groundwater levels. Influence of tidal fluctuation on groundwater levels. | Continuous water levels at site wells and in the Narrows using pressure transducers. Precipitation amounts recorded at area weather stations. |
| Groundwater geochemistry | None. | Location of salt water intrusion and extent of groundwater- surface water interaction. | Groundwater samples will be collected from site wells for field measurements and laboratory analysis of conventional geochemical parameters, salinity. |
| Nature and Extent of Contamination | | | |
| Identify and evaluate source areas | Historical review of Gas Works operations identifies potential source areas. | Potential sources and source areas have not been sufficiently investigated. Potential locations of some potential sources (e.g., tar pits, transfer piping) are unknown or roughly estimated. Lateral and vertical extent of sources in the subsurface is unknown. | Geophysical surveys to identify potential subsurface features. Advance soil borings and complete test pits or trenches in and around potential source areas including former process and residuals management areas: tar pit, residue citern, tar wells; in the ravine fill area; and at a geophysical anomalies detected during geophysical surveys indicating a potential source. Visually observe and record soil stratigraphy and indications of contamination. |
| Evaluate COPCs to determine COCs | Surface and subsurface soil and groundwater samples collected in 2007 and 2008 were analyzed for metals, petroleum hydrocarbons, SVOCs, VOCs and PCBs. | Presence of COPCs previously not evaluated (e.g., cyanide). COPC analysis of sources and in areas not previously characterized. | Samples during the intial investigation phases will be analyzed for all of the COPCs. Soil and groundwater samples collected throughout the investigation will be used in the risk assessment to identify COCs for evaluation of remedial options in the Feasibility Study. |
| Define nature and extent of COCs in soil | Soil samples collected in 2007 and 2008 identified concentrations of metals, PAHs, and VOCs exceeding PRGs. | Current nature and extent of COCs in soil. Presence, nature and extent of COPCs previously not evaluated. | Soil samples will be collected from soil borings, test pits or trenches in and downgradient of source areas, and soil samples using ISM will be collected from soil borings outside source areas, to establish horizontal and vertical limits to the extent of comtamination. Soils will be submitted for chemical analysis of COPCs. |

Table 5-1 - Summary of Existing Information and Data Gaps - UplandsBremerton Gas Works Superfund Site Bremerton, Washington

| Remdial Investigation/Feasbility Study Information Needs by Topic | Existing Information | Data Gaps | Recommended Data Collection | | |
|---|---|--|--|--|--|
| Define nature and extent of COCs in groundwater | Groundwater samples collected in 2007 and 2008 identified concentrations of metals, SVOCs, and VOCs exceeding PRGs. | Current nature and extent of COCs in groundwater Seasonal variability of COCs in groundwater. | Install monitoring wells to evaluate impacts in source areas and establish horizontal and vertical limits to the extent of contamination. Groundwater samples will be initially be collected from all monitoring wells for chemical analysis of all COPCs. A quarterly sampling program will be determined based on initial results. | | |
| Define nature and extent of NAPL | Previous investigations have indicated that NAPL may be present. | | Advance soil borings and complete test pits or trenches in former Gas Works operations and residuals management areas, including the tar pit, residue cistern, tar wells, and in the ravine fill area. Visually observe and record soil stratigraphy and NAPL occurrences. Include monitoring wells screened appropriately to monitor LNAPL (across water table) and DNAPL (above aquitards). Monitor wells for LNAPLs and DNAPL presence. Submit samples of soil and/or NAPL collected from soil borings, test pits, trenches or wells for chemical analysis to characterize NAPL chemistry. If NAPL is identified to be present: advance additional soil borings for deeper NAPL occurences and test pits or trenches for shallow NAPL occurences in areas requiring more precise definition of NAPL occurrences. | | |
| Evaluate potential for recontamination from other area sites | Soil and groundwater samples have been collected from borings and wells located upgradient of the Gas Works property show potential impacts in groundwater south of the property. Limited available data do not show impacts from bulk fuel facilities east of Pennsylvania Avenue or west of Thompson Drive extending onto the Gas Works Property. | Potential impact from adjacent bulk fuel facilities and upgradient industrial sites. | Soil and groundwater data collected from soil borings, test pits and monitoring wells upgradient of the former Gas Works property will be compared to evaluate the extent of contaminants exceeding screening criteria that are associated with the Gas Works site and potential contributions from other area contaminant sources. | | |
| Contaminant Fate and Transport | | | | | |
| NAPL migration pathways | NAPL may be present in the subsurface. MGP-related products include both LNAPL and DNAPL. | Nature and extent of NAPL (see above) NAPL mobility, including NAPL physical characteristics and soil lithology/physical properties | Characterize soil characteristics, NAPL characteristics and extent (see above). Recovery testing to evaluate potential mobility, if NAPL observed in monitoring wells. | | |
| Soil-to-groundwater pathway | Concentrations of Gas Works-associated constituents have been detected above soil and groundwater PRGs. | Leaching potential from contaminated soils. | Include TOC in soil testing program. Collect data to evaluate chemical and geochemical groundwater quality along groundwater transects, located parallel to the groundwater flow direction, to evaluate groundwater conditions with distance from sources. | | |

Table 5-1 - Summary of Existing Information and Data Gaps - Uplands

Bremerton Gas Works Superfund Site Bremerton, Washington

| Remdial Investigation/Feasbility Study Information Needs by | Existing Information | Data Gaps | Recommended Data Collection |
|---|---|--|---|
| Topic | | | |
| Soil-to-surface water pathway | Concentrations of Gas Works-associated constituents have been detected above soil PRGs. | Discharge of contamination through stormwater runoff. | Characterize contamination in sediment and surface water near outfalls. |
| Groundwater-to-surface water pathway | Concentrations of Gas Works-associated constituents have been detected in groundwater above surface water PRGs. | Groundwater transport parameters (velocity, pathway). Attenuation parameters. | Include natural attenuation parameters in groundwater testing program. Characterize hydrogeology and chemical nature and extent (see above). Data may be incorporated into hydrogeologic and fate and transport models. Groundwater monitoring program to assess seasonal variability and long-term trends. |
| Soil-to-air and groundwater-to-air pathway | Concentrations of Gas Works-associated constituents have been detected above current soil and groundwater PRGs. | Potential impacts to future indoor air. | Soil and groundwater data to be used with vapor transport modeling. |
| Human Health and Ecological Risk Assessment | - | | |
| Assess potential receptors and exposure pathways | Concentrations of Gas Works-associated constituents have been detected above current soil and groundwater criteria. | Exposure point concentrations and potential risk to human health through direct contact with soil, ingestion of groundwater, and inhalation via vapor intrusion. Exposure point concentrations and potential risk to ecological receptors through direct contact with soil. | Soil and groundwater chemical analytical results will be compared to human health and ecological risk-based criteria. |

Notes:

BTEX = benzene, toluene, ethylbenzene and xylenes

COC = chemical of concern

COPC = chemical of potential concern

Cs-137 = Cesium 137 isotope

CSL = Cleanup Screening Level

CSO = combined sewer overflow

ISM = incremental sampling methodology

LNAPL = light non-aqueous phase liquide

MGP = manufactured gas plant

NAPL = non-aqueous phase liquid

NOAA: National Oceanic and Atmospheric Administration

PAHs = Polycyclic aromatic hydrocarbons

PRG = preliminary remediation goal

SCO = Sediment Cleanup Objective

SMS = Washington Sediment Management Standards regulations (WAC-173-204)

TOC = total organic carbon

VOC = volatile organic compound

Table 5-2 - Summary of Existing Information and Data Gaps - SedimentsBremerton Gas Works Superfund Site Bremerton, Washington

| RI/FS Information Needs by Topic (What We Need to Know) | Existing Information (What We Already Know) | Data Gaps (What We Don't Know) | Recommended Data Collection (RI Work to Fill Data Gaps) |
|---|---|---|---|
| Nature and Extent of Contamination Assess presence of chemical contaminants associated with historical MGP operations | MGP operational history is well documented. MGP-associated contaminants typically include PAH compounds, selected VOCs (i.e., BTEX compounds), cyanide and dibenzofuran. Surface sediment PAH concentrations within the intertidal beach areas have been extensively sampled. Some testing for other parameters (semivolatiles, metals and VOCs) has also been performed on a more limited basis. | Sampling has not yet been performed in areas offshore of the former MGP dock. Testing has not been performed for cyanide in sediments. Testing for alkylated PAHs has not been performed (these parameters are useful in discriminating PAH sources in sediments). | Collect surface sediment samples from MGP dock area Analyze sediment samples for Site COPCs and alkylated PAH to document the "fingerprint" of MGP-associated PAHs |
| Identify chemical contaminants potentially associated with other historical activities within the Site | include ravine fill, oil handling, CSO/stormwater discharges, adjacent marina operations and miscellaneous industrial | Sampling near non-MGP sources is not sufficient to finalize list of COPCs Testing has not yet been performed offshore of former Sesko Oil dock Testing for alkylated PAHs has not been performed (these parameters are useful in discriminating PAH sources in sediments) | Collect surface sediment samples from former Sesko dock area Analyze sediment samples for Site COPCs and alkylated PAH to document the "fingerprint" of MGP-associated PAHs |
| Define the lateral extent of Site COPCs in surface sediment, including the boundary between Site-associated contamination, and contamination from regional inputs | Surface sediment PAH concentrations within the intertidal beach areas have been extensively sampled. Some testing for other parameters (semivolatiles, metals and VOCs) has also been performed on a more limited basis. Extensive data are available documenting regional sediment quality within Port Washington Narrows and Dyes Inlet. Those data indicate an elevated regional PAH concentrations and the presence of certain other regional contaminants. | The lateral extent of Site-associated PAH contamination has not been determined within Port Washington Narrows. Given the presence of elevated PAH concentrations in regional sediments, additional sampling and "fingerprint" data will be needed to define the boundary between Site-associated PAH contamination and PAH contamination from regional inputs. If other Site COPCs are confirmed, then the lateral extent of these COPCs in surface sediments will need to be determined, including the boundary between Site-associated contamination and contamination from regional inputs. | Collect surface sediment samples from across the initial study area and analyze for Site COPCs |
| Define the vertical extent of Site COPCs in sub-surface sediment, including the potential presence of subsurface hydrocarbon deposits (i.e., sheen or NAPL) | Limited subsurface testing has been performed in the western portion of the intertidal beach to evaluate the vertical extent of PAH contamination and hydrocarbon sheen in that area. Results demonstrated that sediment contamination levels decreased rapidly with depth, and the area containing subsurface hydrocarbon sheen was very limited. | Subsurface testing has not been performed in other areas of the beach. The depth of contamination is therefore not defined in those areas. No surface or subsurface testing has been performed areas offshore of the former MGP dock. Core sampling data are not yet sufficient to assess whether subsurface hydrocarbon deposits (sheen or NAPL) may be present in subsurface sediments other than in the western beach area. | Conduct sediment core sampling and chemical analysis within the initial study area to assess the vertical extent of Site COPCs Include sufficient core sampling locations in nearshore and offshore areas to assess the potential presence of susurface hydrocarbon deposits (sheen or NAPL). |

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Table 5-2 - Summary of Existing Information and Data Gaps - SedimentsBremerton Gas Works Superfund Site Bremerton, Washington

| RI/FS Information Needs by Topic (What We Need to Know) | Existing Information (What We Already Know) | Data Gaps (What We Don't Know) | Recommended Data Collection (RI Work to Fill Data Gaps) |
|---|--|---|--|
| Human Health & Ecological Risk Assessment | | | |
| Assess the site-specific partitioning behavior of PAHs in sediments | Literature data can be used to estimate potential partitioning of PAH compounds between sediment and pore-water. However, these methods may not capture site-specific factors. | No site-specific pore-water testing has been performed to assess PAH partitioning behavior in sediments | Conduct paired analysis of bulk sediment and pore-water PAH concentrations in selected study areas for analysis of site-specific partitioning behavior. |
| Assess potential impacts of Site COPCs to benthic receptors | The potential for benthic impacts can be assessed using bulk sediment chemistry (to be defined as described above) along with toxicity threshold values such as the SMS SCO and CSL values, and/or the EPA narcosis toxicity model. Pore-water PAH data may be used directly to assess potential benthic toxicity using the EPA narcosis toxicity model. | Site-specific bioassay testing could be used along-side bulk sediment chemistry and pore-water testing data to assess potential benthic impacts. The need for bioassay testing can be assessed after review of bulk sediment chemistry and pore-water PAH data to be collected as described above. | Contingent Activity: If applicable, based on review of bulk sediment chemistry and pore-water testing data, collect sediment samples from selected areas for confirmational bioassay testing. This testing could be used to verify predicted impacts and refine the lateral extent of those impacts. |
| Assess potential for Site-associated sediment contaminants to accumulate in the tissues of aquatic organisms | Literature data can be used to estimate potential uptake of PAH or other contaminants in the tissues of aquatic organisms. Reliance on literature data may not capture site-specific factors. | No site-specific tissue testing data or bioaccumulation testing data has been performed | Develop estimates of tissue concentrations based on bulk sediment and pore-water testing data and literature-based biotasediment accmulation factors. Contingent Activity: Where warranted, use tissue testing (preferred) or laboratory bioaccumulation testing (alternate) to directly assess the potential accumulation of Site COPCs in selected aquatic organisms. |
| Document the types and quantities of aquatic species present in the vicinity of the Site | Previous habitat and fish/shellfish resource surveys have been performed in the Port Washington Narrows and Dyes inlet areas, documenting locally-abundant fish and shellfish species. Information regarding current and proposed shellfish growing areas, and historical patterns of fishing and shellfish harvesting are available through state and tribal agencies. Patterns of tribal seafood consumption have been identified in previous surveys of the Suquamish, Tulalip and Squaxin nations. | Additional information is required to document the habitat conditions and the types of seafood species present within the ISA and immediate vicinity. | Conduct baseline shellfish surveys of aquatic habitat and fish/shellfish resources within ISA and immediate vicinity |
| Evaluate potential Site-associated water quality impacts as necessary to support exposure assessments in the human health and ecological risk assessments | No surface water data are currently available for the Site. Regional studies have documented anthropogenic surface water contaminant inputs to Port Washington Narrows and Dyes Inlet, including but not limited to stormwater and CSO discharges. Any Site-specific sampling of surface water quality will need to consider potential off-site sources for measured water quality parameters. | Surface water quality for the Site and vicinity are not currently available as required to support the risk assessment data needs. | Analyze surface water samples for Site COPCs. Samples to be collected from both within the initial study area and at selected background stations within Port Washington Narrows east and west of the Site to provide ambient water quality context. |

Table 5-2 - Summary of Existing Information and Data Gaps - Sediments

Bremerton Gas Works Superfund Site Bremerton, Washington

| RI/FS Information Needs by Topic (What We Need to Know) | Existing Information (What We Already Know) | Data Gaps (What We Don't Know) | Recommended Data Collection (RI Work to Fill Data Gaps) |
|--|---|-----------------------------------|---|
| Sediment Stability and Recovery Processes | | | |
| Assess potential near-bottom currents effects on long-term sediment stability and sediment transport processes within the Site and immediate vicinity. | Peak tidal currents within Port Washington Narrows are understood from existing studies (e.g., NOAA tide and current data). Sediment texture and particle size will be defined during surface sediment testing as described above. | | Conduct empirical measurements of near-bottom and mid- channel tidal currents for use in an analysis of sediment stability and evalution of sediment transport proceses. |
| Quantify sedimentation rates to assess the degree of natural recovery that may be anticipated | Geochronology studies have been performed in several areas of Puget Sound, documenting a general pattern of sedimentation. | | Contingent Activity: If warranted, quantify net sedimentation rates near the Site using geochronology test methods (i.e., thinsection cores analyzed with Cs-137 radio-dating). |

Notes:

BTEX = Benzene, toluene, ethylbenzene, and xylenes

COC = Contaminant of Concern

Cs-137 = Cesium 137 isotope

CSL = Cleanup Screening Level

CSO = combined sewer overflow

EPA = U.S. Environmental Protection Agency

MGP = manufactured gas plant

NAPL = nonaqueous product layer

NOAA = National Oceanic and Atmospheric Administration

PAH = polynuclear aromatic hydrocarbon

RI/FS = Remedial Investigation/Feasibility Study

SCO = Sediment Cleanup Objective

SMS = Washington Sediment Management Standards regulations (WAC-173-204)

VOC = volatile organic compound

Table 5-3 - Risk Assessment Plan, Baseline Human Health Risk Evaluation

Bremerton Gas Works Superfund Site

| Planned Risk Assessment Activities | | | | | Parameters to be Refined in Risk Assessment Technical Memo | |
|---|--|---|--------------------------------|---|---|---|
| | Estimation Framework(s) for | Relevant RI Data to be Used | | | Detailed Risk Characterization | |
| Receptor | Exposure Pathway | (media and measurements) | Endpoint | Interpretative Framework | Parameters ¹ | Contingent Testing ² |
| Subsistence consumer of fish/crab | Dietary TDI will be estimated from dietary consumption of fish and crab, incidental sediment, and surface water. TDI estimates will be developed using EPA tribal framework. Finfish to include finfish tissue from relevant species from Suquamish Group D (halibut, sole, flounder, and rockfish). | Concentrations of chemicals in surface sediment (intertidal and subtidal) and surface water. Bulk sediment data will be used along with applicable BSAFs to estimate chemical concentrations in fish and crab tissue. | ELCR or health HQ ³ | ELCR greater than 1 in 1,000,000 or HQ greater than 1 indicates a chemical of potential concern. ³ | Specific data to be used in evaluation Tissue-specific BSAFs and derivation TDI calculation inputs Applicable toxicity and exposure parameters | Collection of site-specific tissue samples may be proposed as an alternative to use of literaturederived BSAFs. |
| Subsistence consumer of shellfish ⁴ | TDI will be estimated from dietary consumption of shellfish (i.e., clams), incidental sediment, and surface water. TDI estimates will be developed using EPA tribal framework. | Concentrations of chemicals in intertidal surface sediment and surface water. Bulk sediment data will be used along with applicable BSAFs to estimate chemical concentrations in shellfish tissue. | ELCR or health HQ ³ | ELCR greater than 1 in 1,000,000 or HQ greater than 1 indicates a chemical of potential concern. ³ | Specific data to be used in evaluation Tissue-specific BSAFs and derivation TDI calculation inputs Applicable toxicity and exposure parameters | Collection of site-specific tissue samples may be proposed as an alternative to use of literaturederived BSAFs. |
| Recreational beach user | TDI will be estimated from dermal contact and incidental ingestion of sediment and surface water. | Concentrations of chemicals in intertidal surface sediment and surface water. | ELCR or health HQ ³ | ELCR greater than 1 in 1,000,000 or HQ greater than 1 indicates a chemical of potential concern. ³ | Specific data to be used in evaluation TDI calculation inputs Applicable toxicity and exposure parameters | None anticipated |
| Construction/excavation worker in beach areas | TDI will be estimated from dermal contact and incidental ingestion of sediment. Exposure from inhalation of fugitive dust/vapor will be estimated using EPA inhalation dosimetry methodology. Typical subsurface construction activities such as digging for building foundations are expected to extend approximately 3 feet in depth | Concentrations of chemicals in intertidal surface and subsurface sediment (0-6 feet below mud-line) and surface water. | ELCR or health HQ ³ | ELCR greater than 1 in 1,000,000 or HQ greater than 1 indicates a chemical of potential concern. ³ | Specific data to be used in evaluation TDI calculation inputs Applicable toxicity and exposure parameters | None anticipated |
| Construction/excavation worker in upland site areas ⁵ | TDI will be estimated from dermal contact and incidental ingestion of soil. Exposure from inhalation of fugitive dust/vapor will be estimated using EPA inhalation dosimetry methodology. | Concentrations of chemicals in site surface and subsurface soils (0-6 feet below ground surface) and soil vapor (as estimated from soil, groundwater, or vapor data). | ELCR or health HQ ³ | ELCR greater than 1 in 1,000,000 or HQ greater than 1 indicates a chemical of potential concern. ³ | Specific data to be used in evaluation TDI calculation inputs Applicable toxicity and exposure parameters | None anticipated |

Table 5-3 - Risk Assessment Plan, Baseline Human Health Risk Evaluation

Bremerton Gas Works Superfund Site

Bremerton, Washington

| | | anned Risk Assessment Activities | | | Parameters to be Refined in Risk | Assessment Technical Memo |
|--|---|--|----------|---|---|---------------------------------|
| Receptor | Estimation Framework(s) for Exposure Pathway | Relevant RI Data to be Used (media and measurements) | Endpoint | Interpretative Framework | Detailed Risk Characterization Parameters 1 | Contingent Testing ² |
| On-site occupational worker ⁵ | TDI will be estimated from dermal contact and incidental ingestion of soil. Exposure from inhalation of fugitive dust/vapor will be estimated using EPA inhalation dosimetry methodology | Concentrations of chemicals in upland surficial soils (0-3 feet below ground surface) and soil vapor (as estimated from soil, groundwater, or vapor data). | | chemical of potential concern. ³ | Specific data to be used in evaluation TDI calculation inputs Applicable toxicity and exposure parameters | None anticipated |
| | TDI will be estimated from dermal contact and incidental ingestion of soil. Exposure from inhalation of fugitive dust/vapor will be estimated using EPA inhalation dosimetry methodology. TDI from consumption of groundwater will be considered pending further evaluation of groundwater beneficial uses. | Concentrations of chemicals in upland soils (0-6 feet below ground surface), groundwater, and soil vapor (as estimated from soil, groundwater, or vapor data). | | chemical of potential concern. ³ | Specific data to be used in evaluation TDI calculation inputs Applicable toxicity and exposure parameters | None anticipated |

Notes:

- 1. The risk assessment technical memorandum will present the toxicity data and risk estimation inputs to be used, and will highlight any proposed adjustments to EPA-defined default parameters.
- 2. If applicable, the Risk Assessment Technical Memorandum will define supplemental data collection to be used to refine risk estimates. If contingent testing is proposed, the detailed testing plan will be documented in a Sampling and Quality Assurance Plan amendment.
- 3. A hazard index will be used to sum HQs for different chemicals with potentially additive effects (i.e., similar toxicological mode of action).
- 4. Shellfish consumption within Port Washington Narrows is currently subject to harvest restrictions. This evaluate site-related risks associated with future harvesting activities should such restrictions be lifted.
- 5. No water supply wells are located on or near the Former Gas Works property and is not relevant for the construction worker or occupational worker scenario. Consumption of groundwater will be retained as a potential pathway for screening under the future on-site residential scenario, pending further evaluation of groundwater beneficial uses.
- 6. The site and vicinity are zoned for industrial uses, and residential use is not applicable to current or reasonably foreseeable uses. However, the risks associated with potential future on-site residential use will be evaluated to understand potential risks, should alternative site uses occur in the future.

BSAF = biota-sediment accumulation factors

ELCR = excess lifetime cancer risk

EPA = U.S. Environmental Protection Agency

HQ = hazard quotient

RI = Remedial Investigation

TDI = Total Daily Intake

Table 5-4 - Risk Assessment Plan, Baseline Terrestrial Ecological Risk Evaluation

Bremterton Gas Works Superfund Site Bremerton, Washington

| Planned Risk Assessment Activities Estimation Framework(s) for Exposure Relevant RI Data to be Used | | | | Parameters to be Refined in Risk Assemble Risk Characterization Parameters | | |
|---|---|---|---|---|---|---------------------------------|
| Receptor | Pathway | (media and measurements) | Endpoint | Interpretative Framework | 1 | Contingent Testing ² |
| vian predator (e.g., obin) | ORNL soil screening benchmarks). | | Probability of reduced survival, growth, and reproduction of terrestrial bird populations. | HQ comparing upper bound soil concentration to screening levels. | Specific data to be used in evaluation Soil screening levels and/or benchmarks Applicable exposure parameters | None anticipated |
| | TDI will be estimated from consumption of | chemical concentrations in upland surface soils (0-6 feet) cross the site. | | HQ comparing estimated exposures from TDI to TRV based on no-effects and low-effects concentrations. ³ | Specific data to be used in evaluation Toxicity reference values Applicable exposure parameters | None anticipated |
| arnivore (e.g., oyote) | relevant interpretative benchmarks (e.g., | Evaluation of potential current exposure will use chemical concentrations in upland surface soils (0-6 feet) from vegetated areas including the upland embankment and unpaved upland site areas. Evaluation | terrestrial mammal populations. | HQ comparing upper bound soil concentration to screening levels. | Specific data to be used in evaluation Soil screening levels and/or benchmarks Applicable exposure parameters | None anticipated |
| | | f of potential future exposure will use chemical concentrations in upland surface soils (0-6 feet) cross the site. | | HQ comparing estimated exposures from TDI to TRV based on no-effects and low-effects concentrations. ³ | Specific data to be used in evaluation Toxicity reference values Applicable exposure parameters | None anticipated |
| mnivore (e.g., ccoon) | screening levels (Eco SSL) and other relevant interpretative benchmarks (e.g., ORNL soil screening benchmarks). | surface soils (0-6 feet) from vegetated areas including the upland embankment and unpaved upland site areas. Evaluation of potential future exposure will use | | HQ comparing upper bound soil concentration to screening levels. | Specific data to be used in evaluation Soil screening levels and/or benchmarks Applicable exposure parameters | None anticipated |
| | TDI will be estimated from consumption of plants, invertebrates, and incidental ingestion of soil. Plant and invertebrate tissue concentrations estimated using soil-to-tissue regression models. | chemical concentrations in upland surface soils (0-6 feet) cross the site. | | HQ comparing estimated exposures from TDI to TRV based on no-effects and low-effects concentrations. ³ | Specific data to be used in evaluation Toxicity reference values Applicable exposure parameters | None anticipated |
| erbivore (e.g., vole) | interpretative benchmarks (e.g., ORNL soil screening benchmarks). | surface soils (0-6 feet) from vegetated areas including the upland embankment and unpaved upland site areas. Evaluation of potential future exposure will use | Probability of reduced survival, growth, and reproduction of terrestrial mammal populations | HQ comparing upper bound soil concentration to screening levels. | Specific data to be used in evaluation Soil screening levels and/or benchmarks Applicable exposure parameters | None anticipated |
| | TDI will be estimated from consumption of plants and incidental ingestion of soil. Plant tissue concentrations estimated using soil-to-tissue regression models. | chemical concentrations in upland surface soils (0-6 feet) cross the site. | | HQ comparing estimated exposures from TDI to TRV based on no-effects and low-effects concentrations. ³ | Specific data to be used in evaluation Toxicity reference values Applicable exposure parameters | None anticipated |

Table 5-4 - Risk Assessment Plan, Baseline Terrestrial Ecological Risk Evaluation

Bremterton Gas Works Superfund Site Bremerton, Washington

| | Fatimation France, variety for France, va | Planned Risk Assessment Activities Relevant RI Data to be Used | 5 | | Parameters to be Refined in Risk Asses | |
|-----------------------------|--|--|---|---|---|---------------------------------|
| Receptor | Estimation Framework(s) for Exposure Pathway | (media and measurements) | Endpoint | Interpretative Framework | 1 | Contingent Testing ² |
| nsectivore (e.g., shrew) | Soil chemical concentrations will be compared to Eco SSL and other relevant interpretative benchmarks (e.g., ORNL soil screening benchmarks). | Evaluation of potential current exposure will use chemical concentrations in upland surface soils (0-6 feet) from vegetated areas including the upland embankment and unpaved upland site areas. Evaluation of potential future exposure will use | Probability of reduced survival, growth, and reproduction of terrestrial mammal populations | HQ comparing upper bound soil concentration to screening levels. | Specific data to be used in evaluation Soil screening levels and/or benchmarks Applicable exposure parameters | None anticipated |
| | TDI will be estimated from consumption of invertebrates and incidental ingestion of soil. Invertebrate tissue concentrations estimated using soil-to-tissue regression models. | chemical concentrations in upland surface soils (0-6 feet) cross the site. | | HQ comparing estimated exposures from TDI to TRV based on no-effects and low-effects concentrations. ³ | Specific data to be used in evaluation Toxicity reference values Applicable exposure parameters | None anticipated |
| Soil invertebrate | Soil chemical concentrations will be compared to Eco SSL and other relevant interpretative benchmarks (e.g., ORNL soil screening benchmarks). | Evaluation of potential current exposure will use chemical concentrations in upland surface soils (0-6 feet) from vegetated areas including the upland embankment and unpaved upland site areas. Evaluation of potential future exposure will use | Probability of reduced survival, growth, and reproduction of soil invertebrate communities. | HQ comparing upper bound soil concentration to screening levels. | Specific data to be used in evaluation Soil screening levels and/or benchmarks Applicable exposure parameters | None anticipated |
| | Invertebrate tissue concentrations will be estimated using soil-to-tissue regression models. | chemical concentrations in upland surface soils (0-6 feet) cross the site. | | HQ comparing estimated tissue concentrations to TRV based on noeffects and low-effects concentrations. ³ | Toxicity reference valuesApplicable exposure parameters | None anticipated |
| Jpland vegetation | Soil concentrations will be compared to Eco SSL and other relevant interpretative benchmarks (e.g., ORNL soil screening benchmarks). | Evaluation of potential current exposure will use chemical concentrations in upland surface soils (0-6 feet) from vegetated areas including the upland embankment and unpaved upland site areas. Evaluation of potential future exposure will use chemical concentrations in upland surface soils (0-6 feet) cross the site. | Probability of reduced survival, growth, and reproduction plant communities. | HQ comparing upper bound soil concentration to screening levels. | Specific data to be used in evaluation Soil screening levels and/or benchmarks Applicable exposure parameters | None anticipated |

Notes:

2/28/2017

- 1. The risk assessment technical memorandum will present the toxicity data and risk estimation inputs to be used, and will highlight any proposed adjustments to EPA-defined default parameters.
- 2. If applicable, the Risk Assessment Technical Memorandum will define supplemental data collection to be used to refine risk estimates. If contingent testing is proposed, the detailed testing plan will be documented in a Sampling and Quality Assurance Plan amendment
- 3. A hazard index will be used to sum HQs for different chemicals with potentially additive effects (i.e., similar toxicological mode of action).

Eco SSL = ecological soil screening levels

HQ = hazard quotient

ORNL = Oak Ridge National Laboratory

Table 5-5 - Risk Assessment Plan, Baseline Aquatic Ecological Risk Evaluation Bremerton Gas Works Superfund Site Bremerton, Washington

| | | Planned Risk Assessment Activi | ties | | | k Assessment Technical Memo |
|---|---|---|--|--|--|--|
| Receptor | Estimation Framework(s) for Exposure Pathway | Relevant RI Data to be Used (media and measurements) | Endpoint | Interpretative Framework | Detailed Risk Characterization Parameters 1 | Contingent Testing ² |
| Piscivorous mammal (e.g., harbor seal) | Dietary TDI will be estimated from consumption of fish and invertebrates. | Concentrations of chemicals in surface sediment (intertidal and subtidal). Bulk sediment data will be used along with applicable BSAFs to estimate chemical concentrations in fish and invertebrate tissue. | Probability of reduced survival, growth, and reproduction of piscivorous mammal populations. | HQ is ratio of TDI to weight- adjusted mammalian TRV based on low- and no-effects concentrations. | Specific data to be used in evaluation Tissue-specific BSAFs and derivation TDI calculation inputs Exposure parameters Toxicity reference values | Collection of site-specific tissue samples of prey species may be proposed as an alternative to use of literature-derived BSAFs for estimation of dietary TDI. |
| Piscivorous raptor (e.g., osprey) | Dietary TDI will be estimated from consumption of fish. | Concentrations of chemicals in surface sediment (intertidal and subtidal). Bulk sediment data will be used along with applicable BSAFs to estimate chemical concentrations in fish tissue. | Probability of reduced survival, growth, and reproduction of aquatic-dependent bird populations. | HQ is ratio of TDI to avian TRV based on low- and no-effects concentrations. ³ | Specific data to be used in evaluation Tissue-specific BSAFs and derivation TDI calculation inputs Exposure parameters Toxicity reference values | Collection of site-specific tissue samples of prey species may be proposed as an alternative to use of literature-derived BSAFs for estimation of dietary TDI. |
| Shore bird (heron) | Dietary TDI will be estimated from consumption of fish, invertebrates, and incidental consumption of intertidal sediment. | | Probability of reduced survival, growth, and reproduction of aquatic-dependent bird populations. | HQ is ratio of TDI to avian TRV based on low- and no-effects concentrations. ³ | Specific data to be used in evaluation Tissue-specific BSAFs and derivation TDI calculation inputs Exposure parameters Toxicity reference values | Collection of site-specific tissue samples of prey species may be proposed as an alternative to use of literature-derived BSAFs for estimation of dietary TDI. |
| Shore bird (sandpiper) | Dietary TDI will be estimated from consumption of invertebrates and incidental consumption of intertidal sediment. | Concentrations of chemicals in intertidal surface sediment. Bulk sediment data will be used along with applicable biota-sediment accumulation factors (BSAFs) to estimate chemical concentrations in invertebrate tissue. | Probability of reduced survival, growth, and reproduction of aquatic-dependent bird populations. | HQ is ratio of TDI to avian TRV based on low- and no-effects concentrations. ³ | Specific data to be used in evaluation Tissue-specific BSAFs and derivation TDI calculation inputs Exposure parameters Toxicity reference values | Collection of site-specific tissue samples of prey species may be proposed as an alternative to use of literature-derived BSAFs for estimation of dietary TDI. |
| Piscivorous fish (e.g., rockfish) | Surface water chemical concentrations compared directly to AWQC. | Concentrations of chemicals in surface sediment (intertidal and subtidal) and surface water. | Probability of reduced survival, growth, and reproduction of fish populations. | HQ is the ratio of the concentration in surface water to the protective criteria. | Specific data to be used in evaluationAWQC values | None anticipated |
| | Surface water chemical concentrations evaluated using TU calculations for 34 PAHs. | | | HQ is the ratio of the concentration in surface water to the protective criteria. | Specific data to be used in evaluationTU calculations | None anticipated |
| | Fish tissue chemical concentrations will be estimated based on sediment BSAF model compared to tissuebased TRVs. | | | HQ is ratio of tissue burden to tissue based TRV based on lowand no-effects concentrations. ³ | Specific data to be used in evaluation Tissue-specific BSAFs and derivation Exposure parameters Toxicity reference values | Collection of site-specific tissue samples may be proposed as an alternative to use of literature-derived BSAFs for this receptor. |

Table 5-5 - Risk Assessment Plan, Baseline Aquatic Ecological Risk Evaluation

Bremerton Gas Works Superfund Site Bremerton, Washington

| | | Planned Risk Assessment Activ | ities | | Parameters to be Refined in Risl | Assessment Technical Memo |
|---|--|--|--|---|--|--|
| Receptor | Estimation Framework(s) for Exposure Pathway | Relevant RI Data to be Used (media and measurements) | Endpoint | Interpretative Framework | Detailed Risk Characterization Parameters ¹ | Contingent Testing ² |
| Omnivorous fish (e.g., sculpin) | Surface water chemical concentrations will be compared directly to AWQC. | Concentrations of chemicals in surface sediment (intertidal and subtidal) and surface water. | Probability of reduced survival, growth, and reproduction of fish populations. | HQ is the ratio of the concentration in surface water to the protective criteria. | Specific data to be used in evaluation AWQC values | None anticipated |
| | Surface water chemical concentrations will be evaluated using TU calculations for 34 PAHs. | | | HQ is the ratio of the concentration in surface water to the protective criteria. | Specific data to be used in evaluation TU calculations | None anticipated |
| | Fish tissue chemical concentrations will be estimated based on sediment BSAF model compared to tissue-based TRVs. | | | HQ is ratio of tissue burden to tissue based TRV based on lowand no-effects concentrations. ³ | Specific data to be used in evaluation Tissue-specific BSAFs and derivation Exposure parameters Toxicity reference values | Collection of site-specific tissue samples may be proposed as an alternative to use of literature-derived BSAFs for this receptor. |
| Benthivorous fish (e.g., flatfish) | Surface water chemical concentrations will be compared directly to AWQC. | Concentrations of chemicals in surface sediment (intertidal and subtidal) and surface water. | Probability of reduced survival, growth, and reproduction of fish populations. | HQ is the ratio of the concentration in surface water to the protective criteria. | Specific data to be used in evaluation AWQC values | None anticipated |
| | Surface water chemical concentrations will be evaluated using TU calculations for 34 PAHs. | | | HQ is the ratio of the concentration in surface water to the protective criteria. | Specific data to be used in evaluation TU calculations | None anticipated |
| | Fish tissue chemical concentrations will be estimated based on sediment BSAF model compared to tissue-based TRVs. | | | HQ is ratio of estimated tissue concentrations and tissue-based TRV based on low- and no-effects concentrations. ³ | Specific data to be used in evaluation BSAFs and derivation Exposure parameters Toxicity reference values | Collection of site-specific tissue samples may be proposed as an alternative to use of literature-derived BSAFs for this receptor. |
| Benthivorous shellfish (e.g., crab) | Surface water chemical concentrations will be compared directly to AWQC, including the PAH FCVs. | Concentrations of chemicals in surface sediment (intertidal and subtidal) and surface water. | Probability of reduced survival, growth, and reproduction of shellfish populations. | HQ is the ratio of the concentration in surface water to the protective criteria | Specific data to be used in evaluation AWQC values | None anticipated |
| | Surface water chemical concentrations will be evaluated using TU calculations for 34 PAHs. | | | HQ is the ratio of the concentration in surface water to the protective criteria | Specific data to be used in evaluation TU calculations | None anticipated |
| Benthic invertebrates (e.g., benthic infauna community) | Sediment chemical concentrations will be compared to SMS criteria for protection of benthic receptors. | Concentrations of chemicals in surface sediment (intertidal and subtidal) and porewater. | Probability of reduced survival, growth, and reproduction of benthic invertebrate communities. | SMS criteria include the sediment cleanup objective and the cleanup screening level. | Specific data to be used in evaluation SMS numeric values | Site-specific sediment bioassays may be proposed as an alternative to use of numeric SMS criteria. |
| | Bulk sediment chemistry and total organic carbon content will be used along with literature-derived equilibrium partitioning coefficients to estimate sediment porewater concentrations for PAH compounds. | | | Estimated sediment porewater concentrations for 34 PAH compounds will be evaluated using the TU method. | Specific data to be used in evaluation Equilibrium partitioning coefficients Toxic unit calculations | Site-specific sediment porewater collection and analysis may be proposed as an alternative to use of porewater concentration estimates derived from equilibrium partitioning coefficients. |

Table 5-5 - Risk Assessment Plan, Baseline Aquatic Ecological Risk Evaluation

Bremerton Gas Works Superfund Site Bremerton, Washington

| | Planned Risk Assessment Activities | | | | Parameters to be Refined in Risk | Assessment Technical Memo |
|----------|--|--|----------|---|--|--|
| | Estimation Framework(s) for | Relevant RI Data to be Used | | | Detailed Risk Characterization | 2 |
| Receptor | Exposure Pathway | (media and measurements) | Endpoint | Interpretative Framework | Parameters ¹ | Contingent Testing ² |
| ,, | Surface water chemical concentrations will be compared directly to AWQC, including the PAH FCVs. | Concentrations of chemicals in surface sediment (intertidal and subtidal) and porewater. | 1 | HQ is the ratio of the concentration in surface water to the protective criteria. | Specific data to be used in evaluation AWQC values | None anticipated |
| | Bulk sediment chemistry and total organic carbon content will be used along with literature-derived equilibrium partitioning coefficients to estimate sediment porewater concentrations for PAH compounds. | | | Estimated sediment porewater concentrations for 34 PAH compounds will be evaluated using the TU method. | evaluation | Site-specific sediment porewater collection and analysis may be proposed as an alternative to use of porewater concentration estimates derived from equilibrium partitioning coefficients. |

Notes:

- 1. The risk assessment technical memorandum will present the toxicity data and risk estimation inputs to be used, and will highlight any proposed adjustments to EPA-defined default parameters.
- 2. If applicable, the Risk Assessment Technical Memorandum will define supplemental data collection to be used to refine risk estimates. If contingent testing is proposed, the detailed testing plan will be documented in a Sampling and Quality Assurance Plan amendment.
- 3. A hazard index will be used to sum HQs for different chemicals with potentially additive effects (i.e., similar toxicological mode of action).

AWQC = ambient water quality criteria

BSAF = biota-sediment accumulation factors

FCV = final chronic value

HQ = hazard quotient

PAH = polycyclic aromatic hydrocarbon

Table 5-6 – Data Quality Objectives Contamination in Soil and Groundwater

Bremerton Gas Works Superfund Site Bremerton, Washington

| Step | Description | | | |
|------------------------------------|---|--|--|--|
| State the Problem | Additional information is necessary determine the lateral and vertical extent of contamination in soil and groundwater, and evaluate risks to human and ecological receptors. | | | |
| Identify the Goal of the Study | The goals are to: Determine the nature and extent of contaminant concentrations in soil and groundwater exceeding PRGs at the Site. Determine seasonal variability in contaminant concentrations in groundwater. Obtain adequate and representative data from soil and groundwater for use in the Baseline Human Health and Ecological Risk Assessments. Determine the potential for recontamination of the Site from groundwater flowing from adjacent sites. | | | |
| Identify Information Inputs | Information inputs include: Preliminary conceptual site model (CSM) (see Section 4.0). ARARS, RAOS, and PRGS (see Section 3.1). Concentrations of COPCs, including VOC, SVOC, pesticide, PCB, cyanide, dioxin/furan, and metals, in soil. Concentrations of COPCs in groundwater. Site geology and hydrogeology, including groundwater occurrence and flow characteristics. | | | |
| Define the Boundaries of the Study | Spatial boundaries: The horizontal extent of the study area is defined by the upland portion of the ISA. The vertical extent of the study area will be based on bounding contamination (as determined by comparison of analytical data to PRGs) during the course of the study. Based on data collected during the study, the boundaries of the study area will be adjusted as needed to encompass the extent of where contamination from the Site has come to be located. Temporal boundaries: Data of sufficient quality (see Section 3.6.2) from previous investigations (beginning in 2007) to those collected as part of this study will be used. Constraints on data collection: The field work and evaluation of data will be phased in order to allow for refinement to the scopes of work for subsequent RI activities. Other constraints may include limitations due to sampling methods, drilling refusal, encountering subsurface structures (such as piping or foundations), or issues with sampling adjacent properties. | | | |

Table 5-6 – Data Quality Objectives Contamination in Soil and Groundwater

Bremerton Gas Works Superfund Site Bremerton, Washington

| Step | Description |
|---|---|
| Develop the Analytic Approach | Nature and extent of contamination: Analyte concentrations from soil and groundwater samples will be used to determine the study boundaries (defined as the extent of contamination). Sample-specific concentrations will be compared to PRGs (see Section 3.1.3). Data will be evaluated and displayed using figures and tables, and the findings will be used to update the CSM. Risk assessment: Soil and groundwater data will be used to estimate exposure-point concentrations for use in estimating risks based on exposure to soil and groundwater (details will be developed and documented in the Risk Assessment Technical Memorandum). |
| Specify Performance or Acceptance Criteria | Ensure, through data review and validation, that the analytical data for collected samples are within acceptable quality limits as defined by applicable EPA data quality protocols (Appendix A, Upland SQAPP). |
| | Ensure that sampling and analytical representativeness allow for adequate delineation of contaminant nature and extent, and estimates of exposure for the risk assessment, and subsequent identification of areas and media requiring remediation. |
| Develop the Plan for Obtaining Data | The detailed plan for obtaining data is presented in this work plan and accompanying Upland SQAPP (Appendix A). A stepwise approach is proposed to determine the extent of contamination in soil and groundwater: |
| | Investigate and identify potential sources through geophysical surveys Investigate potential sources via trenches, test pits, and borings at locations of historical Site features and subsurface anomalies identified by the geophysical surveys. Delineate sources based on field observations. Collect samples of source materials to evaluate the types of contaminants associated with each. Collect samples of soil beneath potential sources to evaluate vertical extent of contaminants. Analyze samples for all COPCs. |
| | Characterize soil immediately downgradient of source areas with deep borings, collecting samples of fill, vadose zone, shallow water table, deep water table/aquitard, and lower aquifer soils if present. Analyze samples for all COPCs. Determine depths and locations of wells in and downgradient of Source Areas based on soil data, install wells, and characterize groundwater. Evaluate contaminant concentrations in soil outside source areas using |
| | incremental sampling methodology (ISM) to a depth of 6 feet. Install monitoring wells outside source areas and analyze groundwater for all COPCs to determine the extent of contamination above PRGs. |
| | Conduct quarterly monitoring of contaminants in groundwater at monitoring wells for a minimum of 1 year to assess seasonal variability. |

Table 5-6

Table 5-7 – Data Quality Objectives Sources of Contamination (Upland) Bremerton Gas Works Superfund Site

Bremerton, Washington

| Step | Description |
|---|---|
| State the Problem | Additional information is necessary to identify the location of historical sources of contamination at the Site. |
| Identify the Goal of the Study | The goals are to: Determine locations where contaminants may have been released for the purposes of targeting subsurface investigations. Evaluate the potential presence of subsurface features that may act as a source or conduit of contamination. Delineate the source boundaries and/or estimate the source dimensions. Identify the types of contaminants associated with each source. |
| Identify Information Inputs | Information inputs include: Preliminary conceptual site model (CSM) (see Section 4.0). Contaminant concentrations in soil, groundwater, and source materials (e.g., NAPL). Site geology including fill composition and occurrence. Historical information, including aerial photographs. Utility and geophysical surveys. Subsurface survey through observation of targeted, shallow excavations (borings, test pits, and trenches). |
| Define the Boundaries of the Study | Spatial Boundaries: The horizontal extent of the study area is defined by the extent of historical gas works operations, including the fill areas in the former ravine and along the shoreline. The vertical extent of the study area will be based on bounding the depths of sources, including the depth of fill, during the course of the study as feasible for the exploration tools used. Constraints on data collection: The field work and evaluation of data will be phased in order to allow for evaluation of initial data to inform subsequent RI activities. Other constraints may include limitations due to drilling refusal, stability of trenches/test pits; encountering subsurface features that affect survey equipment response; or access issues with sampling adjacent properties. |
| Develop the Analytic Approach | Collected information, survey data, and observations will be used to identify areas for further exploration and sampling. Analyte concentrations in source materials will be used to evaluate source composition. Data will be evaluated and displayed using figures and tables, and the findings will be used to update the CSM. |
| Specify Performance or Acceptance Criteria | Geophysical surveys are a qualitative evaluation. For analytical sampling of sources: Ensure through data review and validation that the analytical data for collected samples are within acceptable quality limits as defined by applicable EPA data quality protocols (Appendix A, Upland SQAPP). Ensure that sampling and analytical representativeness allow for adequate characterization of different potential sources. |

Table 5-7

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Table 5-7 – Data Quality Objectives Sources of Contamination (Upland) Bremerton Gas Works Superfund Site

Bremerton Gas Works Superfund Site Bremerton, Washington

| Step | Description |
|--|--|
| Develop the Plan for Obtaining Data | The detailed plan for obtaining data is presented in this work plan and accompanying Upland SQAPP (Appendix A). A stepwise approach is proposed to identify potential sources: |
| | Utility/geophysical surveys will be used to update Site maps of potential sources and target explorations. |
| | Historical and survey data will be used to locate explorations (borings, test pits, or trenches). |
| | The lateral and vertical extent of sources will be determined based on field observations. |
| | Representative samples of source materials will be collected from shallow excavations for chemical analysis. |
| | Alignment of buried pipes, if encountered, will be further located as practicable using utility location techniques. |

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Table 5-8 – Data Quality Objectives Site Physical Characteristics Bremerton Gas Works Superfund Site

| Step | Description |
|--|--|
| State the Problem | Additional information is necessary to characterize Site physical characteristics. |
| Identify the Goal of the | The goals are to: |
| Study | Determine soil lithology and physical properties of lithologic units. |
| | Determine hydraulic characteristics of Site aquifer units. |
| | Understand role of tidally-influenced surface water on groundwater flow. |
| | Evaluate whether Site groundwater is a potential drinking water source. |
| Identify Information | Information inputs include: |
| Inputs | Preliminary conceptual site model (CSM) (see Section 4.0). |
| | Logging of Site soil lithology from subsurface explorations. |
| | Physical soil characteristics, including gradation, density, Atterberg limits, penetration tests, and moisture content. |
| | Hydraulic conductivity of aquifer units. |
| | Water levels at Site wells throughout seasonal and tidal cycles. |
| | Salinity data at Site monitoring wells. |
| Define the Boundaries of the Study | The study area is defined by the upland portion of the ISA. The horizontal boundaries of the study area will be adjusted as needed to encompass the extent of where contamination from the Site has come to be located. The vertical extent of the study area will extend to include lithologic and aquifer units to define the vertical extent of contaminants in soil and groundwater at concentrations exceeding the PRGs. |
| Develop the Analytic Approach | Identify distinct lithologic and aquifer units through soil sampling. Submit representative samples from each unit for physical testing (Appendix A). Conduct hydraulic testing of aquifer units at representative monitoring wells (Appendix A). |
| Specify Performance or Acceptance Criteria | Physical data will be collected and analyzed using standard test measurements and procedures. Soil lithology characterization and sampling will be performed under the supervision of a registered geologist. Hydraulic testing will be performed under the supervision of a registered hydrogeologist. |
| Develop the Plan for Obtaining Data | The detailed plan for obtaining data is presented in this work plan and accompanying Upland SQAPP (Appendix A). All subsurface explorations at the Site will be logged, and representative samples from each distinct lithologic unit will be analyzed for physical parameters. An initial study of tidally influenced groundwater flow will be conducted using water-table wells, and a limited number of deeper wells, to develop a preliminary estimate of groundwater flow characteristics and assist in locating subsequent explorations. A subsequent tidal study and hydraulic testing will be performed for contaminated aquifer units after the vertical and lateral limits of contaminated groundwater are determined. |

Table 5-9 – Data Quality Objectives NAPL Characterization

Bremerton Gas Works Superfund Site Bremerton, Washington

| Step | Description |
|---------------------------------------|---|
| State the Problem | Additional information is required to characterize the extent of NAPLs, their physical and chemical characteristics, and their potential mobility. |
| Identify the Goal of the Study | The goals are to: Determine the lateral and vertical boundaries of NAPL occurrences. Characterize soil characteristics surrounding NAPL occurrences. Identify physical and chemical characteristics of NAPL. Evaluate NAPL mobility. |
| Identify Information Inputs | Information inputs include: Preliminary conceptual site model (CSM) (see Section 4.0). Logging of Site soil lithology from subsurface explorations. Field observations of potential NAPL indicators. Chemical concentrations of contaminants in soil samples where NAPL may be observed. Measurements of NAPL presence and thickness in monitoring wells. Analysis of NAPL samples for physical properties, including viscosity, density, and flash point, and chemical composition. |
| Define the Boundaries of the Study | The study area is defined by the upland portion of the ISA. The boundaries of the study area will be adjusted as needed to encompass the extent of where contamination from the Site has come to be located. The vertical extent of the study area will extend to include geologic units to the maximum depth of NAPL extent. Constraints on data collection: The field work and evaluation of data will be phased in order to allow for evaluation of initial data to inform subsequent RI activities. Limitations may include depth limitations on exploration techniques (refusal during drilling or test pit/trench stability), subsurface obstructions such as utilities, surface obstructions such as buildings, and access issues on adjacent properties. Different tools (auger or sonic drilling) may be utilized, as needed, to achieve required depths. |
| Develop the Analytic Approach | Assess subsurface soil during logging for the potential presence of NAPL and to characterize soil lithology around potential NAPL occurrences. Where potential NAPL is observed, collect samples of potential NAPL-impacted soil for chemical analysis. Install monitoring wells at locations of potential NAPL occurrence and gauge for NAPL presence and thickness. Where measurable NAPL is observed in monitoring wells, collect NAPL samples for laboratory analysis. Where NAPL is observed in the subsurface, contingent studies for characterizing the lateral and vertical extent of NAPL include the TarGOST technology (see Section 5.6). Where sufficient NAPL is measured in monitoring wells, contingent studies for characterizing mobility and recoverability of NAPL include baildown testing at representative wells containing NAPL (see Section 5.6). |

Table 5-9

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Table 5-9 – Data Quality Objectives NAPL Characterization

Bremerton Gas Works Superfund Site Bremerton, Washington

| Step | Description |
|------------------------|--|
| Specify Performance or | Physical and chemical testing of NAPL samples to be conducted following EPA- |
| Acceptance Criteria | approved and/or standard test methods. Soil logging to be performed under the |
| | supervision of a registered geologist. |
| Develop the Plan for | The detailed plan for obtaining data is presented in this work plan and accompanying |
| Obtaining Data | Upland SQAPP (Appendix A). NAPL investigations will work from the known to the |
| | unknown, starting with suspected source areas and extending outward from identified |
| | sources until the lateral and vertical extent of NAPL is identified. |

Table 5-10 – Data Quality Objectives Contaminant Fate and Transport Bremerton Gas Works Superfund Site

| Step | Description |
|---|---|
| State the Problem | Additional information is necessary to characterize contaminant transport and attenuation at the Site. |
| Identify the Goal of the Study | The goals are to: Evaluate contaminant transport within and partitioning between environmental media. Evaluate potential mechanisms for contaminant attenuation. |
| Identify Information Inputs | Information inputs include: Logging of Site soil lithology from subsurface explorations. Total organic carbon in soil and sediment. Chemical concentrations of contaminants in soil, groundwater, sediment, and surface water. Conventional geochemical parameters in groundwater, including sodium, calcium, potassium, magnesium, chloride, nitrate, nitrite, sulfate, sulfide, alkalinity, ferrous and ferric iron, dissolved manganese, organic carbon, dissolved oxygen, pH, and temperature. |
| Define the Boundaries of the Study | The study area is defined by the ISA. The boundaries of the study area will be adjusted as needed to encompass the extent of where contamination from the Site has come to be located. |
| Develop the Analytic Approach | Assess subsurface soil lithology to evaluate potential preferential migration pathways. Collect and analyze representative samples of each lithologic unit for total organic carbon for evaluations of leaching and sorption. Qualitatively evaluate geochemical parameters, in conjunction with contaminant data, to assess potential for ongoing natural attenuation of contaminants. Conduct vapor intrusion modeling to assess potential contaminant concentrations in indoor air, if structures were present. If the extent of contamination and modeling results indicate a potential exposure risk, soil vapor and/or indoor air sampling may be conducted. See Section 5.6, contingency studies. |
| Specify Performance or Acceptance Criteria | Ensure thorough data review and validation that the analytical data for collected samples are within acceptable quality limits as defined by applicable EPA data quality protocols. |
| Develop the Plan for Obtaining Data | The detailed plan for obtaining data is presented in this work plan and accompanying Upland SQAPP (Appendix A). Lithologic characterization and collection of samples for total organic carbon analysis will be performed during soil and sediment investigations. Geochemical monitoring will be included in groundwater monitoring program. |

Table 5-11 – Data Quality Objectives—Habitat and Intertidal Shellfish Surveys

Bremerton Gas Works Superfund Site Bremerton, Washington

| Step | Description |
|---|--|
| State the Problem | Additional information is necessary to define intertidal and subtidal baseline habitat conditions within Port Washington Narrows. The information will not be used for risk determination or consumption rates. |
| ldentify the Goals of the Study | Evaluate intertidal and subtidal habitat characteristics within the Site and vicinity, including differences in sediment grain size, vegetation, epifauna, other fish and wildlife and physical features (e.g., bed rock outcropping or anthropogenic features). |
| | Quantify the existing abundance of shellfish resources in beach areas of the initial study area (ISA) to establish baseline conditions. |
| Identify Information | Information inputs include: |
| Inputs | Visual surveys of intertidal and subtidal habitat characteristics within the Site and vicinity, are needed to identify structures and differences in sediment grain size, vegetation, epifauna, and identify habitat for other fish and wildlife. |
| | Direct baseline assessment of the abundance of current shellfish in beach areas in within and the immediate vicinity of the Initial Study Area (ISA). |
| Define the Boundaries of the Study | Visual surveys of intertidal and subtidal habitat characteristics will extend throughout the ISA, including transects located in parallel and perpendicular to the axis of the narrows and located along different depth profiles. |
| | Baseline assessment to be conducted within and the immediate vicinity of the ISA. |
| Develop the Analytic Approach | Visual surveys will be performed using a towed camera with integrated DGPS position logging so that all visual observations may be georeferenced. |
| | The baseline shellfish assessment will be performed using methods developed by the Washington Department of Fish and Wildlife for this purpose (Appendix B). |
| Specify Performance or Acceptance Criteria | The DGPS position logging will be verified during the visual surveys to confirm the accuracy of survey locating. The visual quality of the survey will be monitored during collection with a real-time video feed to verify the usability of collected footage. |
| | Surveys of current shellfish resources will comply with quality assurance/quality control protocols developed by the Washington Department of Fish and Wildlife. Baseline assessment results will not be used to inform risk assessment. |
| Develop the Plan for Obtaining Data | The detailed plan for obtaining data is presented in this work plan and accompanying Marine Sampling and Quality Assurance Project Plan (SQAPP; Appendix B). |
| - | Visual surveys will be conducted by towed camera surveys with position logging and real-time video feed for confirming data acquisition. Planned survey transects are defined in the SQAPP. |
| | Shellfish surveys will be conducted during low-tide events following applicable WDFW methodologies. The planned sampling locations are defined in the SQAPP. |

Data Quality Objectives—Contamination in Surface SedimentBremerton Gas Works Superfund Site

Bremerton, Washington

| Step | Description |
|--|--|
| State the Problem | Additional information is necessary to determine the lateral extent of contamination in intertidal and subtidal surface sediment (0–4-inch depth interval). Based on data collected during the study, the boundaries of the study area will be adjusted as needed to encompass the extent of where contamination from the Site has come to be located. The data will also provide the information necessary to support the evaluation of risks to human health and ecological receptors exposed to surface sediment. |
| Identify the Goals of the Study | Determine the nature and extent of contaminant concentrations in surface sediment, exceeding PRGs at the Site. Obtain adequate and representative data from surface sediment for use in the Baseline Human Health and Ecological Risk Assessment Evaluate Site-specific polycyclic aromatic hydrocarbon (PAH) bioavailability in sediment porewater relative to literature-derived partitioning coefficients. |
| Identify Information Inputs | Information inputs include: Initial sampling in potential source areas for comprehensive physical and chemical testing to identify preliminary contaminants of potential concern and characterize the lateral extent of contamination in surface sediments. This data inputs will also be used to inform the risk assessment process. Sampling for comprehensive physical and chemical parameters in intertidal and subtidal surface sediment within the ISA. Paired sampling of PAHs in bulk sediment and porewater at selected locations to evaluate partitioning. |
| Define the Boundaries of the Study | Chemical testing of surface sediments will be conducted within the Initial Study Area. However, based on data collected during the study, the boundaries of the study area will be adjusted as needed to encompass the extent of where contamination from the Site has come to be located. Temporal boundaries: Data of sufficient quality (see Section 3.6.2) from previous sediment investigations (beginning in 2010) to those collected as part of this study will be used, if necessary to delineate the study boundary Constraints on data collection: The field work and evaluation of data will be phased in order to allow for evaluation of initial data to inform subsequent RI activities. Other limitations may result from sampling methodology such as refusal. |
| Develop the Analytic Approach | Nature & Extent of Contamination: Chemical testing results from sediment samples will be used to determine the study boundaries (defined as the extent of contamination). Sample-specific concentrations will be compared to PRGs (see Section 3.1.2). Data will be evaluated and displayed using figures and tables, and the findings will be used to update the CSM. Risk Assessment: Sediment data will be used to estimate exposure point concentrations for use in estimating risks as part of the risk assessment technical memo). Porewater PAH concentrations will also be evaluated using the EPA (2003) equilibrium partitioning sediment benchmark framework. |

Table 5-12

Data Quality Objectives—Contamination in Surface SedimentBremerton Gas Works Superfund Site

| Step | Description |
|--|---|
| Specify Performance or Acceptance Criteria | Ensure thorough data review and validation that the analytical data for collected samples are within acceptable quality limits as defined by applicable EPA data quality protocols (Appendix B, Marine SQAPP). |
| | Ensure that sampling and analytical representativeness allow for adequate delineation of contaminant nature and extent and estimates of exposure for the risk assessment, and subsequent identification of areas and media requiring remediation. |
| Develop the Plan for Obtaining Data | The detailed plan for obtaining data is presented in this work plan and accompanying Marine SQAPP. |
| | Initial sampling locations were identified during RI/FS scoping and discussions with the EPA project team based on historical source areas, previous sampling results, and an analysis of potential sediment fate and transport processes. This sampling plan is identified in the SQAPP. |
| | The Administrative Order on Consent (AOC) and RI/FS Work Plan include contingencies for additional sampling, should the nature and extent of Site-related contamination not be fully delineated during the initial sampling effort. |

Data Quality Objectives—Contamination in Subsurface SedimentBremerton Gas Works Superfund Site

Bremerton, Washington

| Step | Description |
|--|--|
| State the Problem | Additional information is necessary to determine the lateral and vertical extent of contamination in intertidal and subtidal subsurface sediment (greater than 4-inch depth interval) Define the Boundaries" step: "Based on data collected during the study, the boundaries of the study area will be adjusted as needed to encompass the extent of where contamination from the Site has come to be located and to provide information necessary to support the evaluation of human health risks for exposures to subsurface sediment in intertidal areas. |
| Identify the Goals of the Study | Determine the nature and extent of contaminant concentrations in subsurface sediment exceeding applicable PRGs. Obtain adequate and representative data from surface sediment for use in the Baseline Human Health and Ecological Risk Assessment. |
| Identify Information Inputs | Information inputs include: Document sediment stratigraphy at each coring location, including screening for potential presence of NAPL, hydrocarbon contamination or other anthropogenic impacts. Quantify concentrations of a comprehensive suite of chemicals in subsurface sediments from a minimum of two subsurface depth intervals, representing the zone of highest apparent contamination and the top of the uncontaminated sediment layer. Analysis of additional archived sediment samples may be required depending on the results of initial sample analysis. |
| Define the Boundaries of the Study | Confirm sediment stratigraphy with selected analysis of sediment grain size. Spatial boundaries: The vertical extent of the study area is defined by contaminants in the subsurface sediments (as determined by comparison of analytical data to PRGs) during the course of the study. Based on data collected during the study, the vertical boundaries of the study area will be adjusted as needed to encompass the extent of where contamination from the Site has come to be located. Temporal boundaries: Data of sufficient quality (see Section 3.6.2) from previous sediment subsurface investigations (beginning in 2013) to those collected as part of this study will be used. Constraints on data collection: The field work and evaluation of data will be phased in order to allow for evaluation of initial data to inform subsequent RI activities Other limitations may result from sampling methodology such as refusal. |
| Develop the Analytic Approach | Vertical Nature & Extent of Contamination: Chemical testing results from subsurface sediment samples will be used to determine the vertical extent of contamination. Sample-specific concentrations will be compared to PRGs (see Section 3.1.2). Data will be evaluated and displayed using figures and tables, and the findings will be used to update the CSM. |
| Specify Performance or Acceptance Criteria | Ensure thorough data review and validation that the analytical data for collected samples are within acceptable quality limits as defined by applicable EPA data quality protocols (Appendix B, Marine SQAPP). Ensure that sampling and analytical representativeness allow for adequate delineation of contaminant nature and extent and subsequent identification of areas and media requiring remediation. |

Table 5-13

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Data Quality Objectives—Contamination in Subsurface SedimentBremerton Gas Works Superfund Site

| Step | Description |
|--|---|
| Develop the Plan for Obtaining Data | The detailed plan for obtaining data is presented in this work plan and accompanying Marine SQAPP. |
| | Initial sampling locations were identified during RI/FS scoping and discussions with the EPA project team based on historical source areas, previous sampling results, and an analysis of potential sediment fate and transport processes. This sampling plan is identified in the SQAPP. |
| | The Administrative Order on Consent (AOC) and RI/FS Work Plan include contingencies for additional sampling, should the nature and extent of Site-related contamination not be fully delineated during the initial sampling effort. |

Data Quality Objectives—Contamination in Surface WaterBremerton Gas Works Superfund Site

Bremerton, Washington

| Step | Description |
|--|---|
| State the Problem | Additional information is necessary to determine the nature and extent of Site-related contaminants surface water and to support the evaluation of human health and ecological risks. |
| Identify the Goals of the Study | Determine the nature and extent of confirmed contaminant concentrations in surface water at the Site. Obtain adequate and representative data from surface sediment for use in the Baseline Human Health and Ecological Risk Assessment. |
| Identify Information Inputs | Information inputs include: Measurement of a comprehensive suite of chemicals in surface water at locations within the ISA and at background stations within Port Washington Narrows. Parallel testing for conventional parameters, including total organic carbon, dissolved organic carbon, total suspended solids, dissolved oxygen, pH, salinity, and temperature. |
| Define the Boundaries of the Study | Spatial boundaries: The extent of the study area is defined by Site-related comprehensive chemical testing in the surface water (as determined by comparison of ISA results background station and applicable water quality criteria) during the course of the study. Based on data collected during the study, the boundaries of the study area will be adjusted as needed to encompass the extent of where contamination from the Site has come to be located. |
| Develop the Analytic Approach | Nature & Extent of Contamination: Chemical testing results will be used to determine the extent of contamination in surface water. Sample-specific concentrations will be compared to background stations and applicable water quality criteria. Data will be evaluated and displayed using figures and tables, and the findings will be used to update the CSM. Temporal boundaries: Data will be collected during in the dry and wet seasons to determine any temporal trends in surface water quality. |
| Specify Performance or Acceptance Criteria | Ensure thorough data review and validation that the analytical data for collected samples are within acceptable quality limits as defined by applicable EPA data quality protocols (Appendix B, Marine SQAPP). Ensure that sampling and analytical representativeness allow for adequate delineation of contaminant nature and extent and estimates of exposure for the risk assessment, and subsequent identification of areas and media requiring remediation. |
| Develop the Plan for Obtaining Data | The detailed plan for obtaining data is presented in this work plan and accompanying Marine SQAPP. Surface water sampling locations include two areas within the ISA that could potentially be impacted by releases from groundwater or sediment. Two background locations within Port Washington Narrows are included to help differentiate potential Site-related impacts and contamination from off-Site sources. Four rounds of sampling are included to assess the potential seasonal variability in surface water contaminant concentrations. |

Table 5-14

Data Quality Objectives—Marine Area Sediment Stability and Recontamination Processes

Bremerton Gas Works Superfund Site Bremerton, Washington

| Step | Description |
|--|---|
| State the Problem | Additional information is necessary to evaluate the stability of existing Site sediments and to support the evaluation of potential recontamination from migration of off-Site sediments. |
| Identify the Goals of the Study | Quantify near-bottom tidal currents within Port Washington Narrows for use, along with sediment grain size distribution quantified in other RI activities, in evaluating the stability of the existing bed sediments within the ISA. |
| | Quantify the physical characteristics of surface sediments in adjacent areas of Port Washington Narrows to support the FS evaluation of assess sediment movement and deposition within the Port Washington Narrows. |
| Identify Information | Information inputs include: |
| Inputs | Measurements of the direction and velocity of tidal currents in mid depth and near-bottom areas of Port Washington Narrows during relatively strong and approximately average ingoing and outgoing tides. |
| | Measurements of the physical parameters of surface sediments in off-Site areas of Port Washington Narrows. |
| | Modeled wind and wave action to supplement measured tidal current data |
| Define the Boundaries of the Study | The boundary for the tidal current study includes four transects extending south to north across Port Washington Narrows extending from the Former Gas Works and adjacent beach areas out beyond the boundaries of the ISA. |
| | The boundary for the study of surface sediment quality within Port Washington Narrows extends from the ISA east and west to the ends of Port Washington Narrows. |
| Develop the Analytic Approach | Tidal currents will be measured using a vessel-mounted Acoustic Doppler Current Profiler in order to document changes in tidal currents with depth, location, and time during the course of a daily tide cycle with strong ingoing and outgoing tides. |
| | Physical measurements (total organic carbon, total solids, and grain size) of intertidal and subtidal surface sediments of Port Washington Narrows will be quantified using EPA-approved methods. Archive samples will be sampled and frozen for potential chemical testing, if necessary, to inform recontamination potential evaluations. |
| Specify Performance or Acceptance Criteria | Ensure thorough data review and validation that the analytical data for collected samples are within acceptable quality limits as defined by applicable EPA data quality protocols (Appendix B, Marine SQAPP). |
| | Ensure that sampling and analytical representativeness allow for adequate characterization sediment physical characteristics within the Port Washington Narrows. |

Data Quality Objectives—Marine Area Sediment Stability and Recontamination Processes

Bremerton Gas Works Superfund Site

Bremerton, Washington

| Step | Description |
|--|--|
| Develop the Plan for Obtaining Data | The detailed plan for obtaining data is presented in this work plan and accompanying Marine SQAPP. |
| | Tidal surveys will be conducted by a qualified contractor along transects at the specified locations during a tidal cycle with strong ingoing and outgoing tides. |
| | Sediment sampling locations were selected to include both areas subject to potential sediment movement by littoral drift and sediments subject to potential current-induced sediment movement. |
| | In order to allow for the use of archived mass to inform recontamination potation, if necessary, sampling locations were adjusted to avoid areas likely to be impacted by known or suspected contaminated sites or potential pollution sources |

Page 1 of 2

Table 5-16 Summary of Marine Sampling Design

Bremerton Gas Works Superfund Site Bremerton, Washington

| Area | Sub-Area | Sample Type | Purposes | Purposes Number of Samples and Location Rationale | | | |
|--------------------------------|--|---|--|--|--|--|--|
| Sediment Sampling | | | | | | | |
| | | | To define the horizontal nature and extent of contamination in intertidal sediments | Bulk chemistry at 5 intertidal stations collected throughout beach area adjacent to former Gas Works and ravine | PAHs (including alkylated), TS, TOC, grain size | | |
| | Co-Located Intertidal | Intertidal Grab Samples | Evaluate concentrations of metals, SVOC and cyanide along Gas Works intertidal area | Supplemental testing for bulk chemistry at 5 intertidal stations adjacent to former Gas Works and ravine | Cyanide (total and available), metals and SVOC | | |
| | and Subtidal Sediment Grabs and | | Evaluate pore-water concentrations of PAH and alkylated PAH concentrations | Pore-Water chemistry at 5 intertidal stations | PAHs (including alkylated) in pore-water | | |
| Initial Study Area | Cores | Subtitdal Grab Samples | To define the horizontal nature and extent of contamination in subtidal sediments | 14 subtidal stations collected in transects down the slope toward to the channel elevation. | PAHs (including alkylated), TS, TOC, grain size | | |
| | | Vibracores | To define the vertical nature and extent of contamination in intertidal and subtidal sediments in including NAPL and Sheens | 5 intertidal and 12 subtidal stations Advanced in transects down the slope toward to the channel elevation and two within the marina. | PAHs, TS, TOC | | |
| | Other Intertidal and Subtidal Sediment Grabs | Intertidal Grabs | Provide bounding to the nature & extent of site-associated impacts in intertidal sediment | 2 stations Step-out sampling in accessible intertidal areas within eastern extent of the ISA. The western intertidal extent is a rip rap armored slope and not generally accessible. | PAHs (including alkylated), TS, TOC, grain size | | |
| | Glabs | Subtidal Grabs | To define the horizontal nature and extent of contamination in subtidal sediments | 12 stations Step-out sampling between slope area and ISA boundary. | PAHs (including alkylated), TS, TOC, grain size | | |
| | Intertidal | Surface Grab al (Multi-Increment Composite) | Document quality of intertidal sediments within Port Washington Narrows to provide an estimate of recontamination potential from sediment movement (littoral drift and bed load) and deposition | 11 stations Collection along north side and five along the south side of the narrows. Stations placed in publically accessible intertidal areas. | PAHs (including alkylated), TS, TOC, grain size | | |
| Port Washington Narrows | intertidal | | Evaluate relationship between predicted and actual pore-water concentrations of PAH and alkylated PAH | 5 stations Representative samples of Narrows intertidal samples (every other sample). Allows estimate of central tendency. | PAHs (including alkylated) | | |
| | Subtidal Surfa | | Document quality of intertidal sediments within Port Washington Narrows to provide an estimate of recontamination potential from sediment movement (sediment bed load) and deposition | 6 stations Collection along the general centerline and deeper sections of the channel. | PAHs (including alkylated), TS, TOC, and Grain Size | | |
| Surface Water Sam | pling | | | | | | |
| Initial Study Area | | Grab | Quantify concentrations of site-associated COPCs in surface water | 2 locations Seasonal sampling at 2 depths per location | Conventional Parameters, PAHs (including alkylated) | | |
| Port Washington Narrows | Surface Water Grab | | Quantify concentrations of COPCs in surface water to assess potential regional influences | 2 locations Seasonal sampling at 2 depths per location | Conventional Parameters, PAHs (including alkylated) | | |
| Habitat and Physica | Habitat and Physical Surveys | | | | | | |
| Initial Study Area and Port | Intertidal | Visual & Photo Survey | Conduct surveys of aquatic habitat and fish/shellfish resources near the Site within Port Washington Narrows. | 5 locations within ISA intertidal area, and 11 locations within Port Washington Narrows | Visual survey for clam identification and abundance | | |
| Washington Narrows | Subtidal | Towed-Camera Survey | Refine environmental setting information | 6 transects perpendicular to and 5 transects in parallel with the Port Washington Narrows | Mapping of substrate, vegetation and identified aquatic species | | |
| Initial study area | Subtidal | ADCP Transects | Measure Near-bottom currents that may impact sediment stability | 4 transects perpendicular to Port Washington Narrows (2 tide conditions) | Conduct empirical measurements of near-bottom and mid- channel tidal currents for use in an analysis of sediment stability. | | |

Notes

1. Samples to be archived frozen for contingent analysis should additional testing be required for SVOC or heavy metals.

PAH = polynuclear aromatic hydrocarbons NA = not applicable

TBD = to be determined

TOC = total organic carbon

ADCP = acoustic doppler current profiler

NAPL = non-aqueous phase liquid COPCs = chemicals of potential concern SVOC = semi-volatile organic compound

TS = total solids

Table 7-1 - Remedial Technologies for NAPL

Bremerton Gas Works Superfund Site Bremerton, Washington

| Deed restrictions addressing soil of distributions are offer as decision and distributions are offer groundwister wells grown and sold restrictions and distributions are offer as decision of the properties. The properties of the | NAPL General Response Actions | Remedial Technology | Process Options | Description |
|--|-------------------------------------|---|---|---|
| monitoring to prevent disturbance of engineered controls Lee Restrictions Dead rentations Dead rentations with other technology approaches. Control lateral movement of NAPL by excavating at trench and baciding with a love-persoaches, distinged from the rentation of NAPL. Dead rentation in the dead rentation and the rentation of NAPL by installing polition of water, variations of the Pull by preparation with the rentation brought spirity species benefits Dead rentation brought spirity spirity species benefits Dead rentation brought spirity spirity s | | | signs to control Site | Signs, fences, or other measures to prevent access to the Site. |
| Situry Well Surry Well Su | Institutional Controls ¹ | Use Restrictions | monitoring to prevent disturbance of | that may interfere with a cleanup action or result in exposure to |
| Surry Wall Control lateral movement of NAPL by installing (driving or vibrating) steel or plastic sheet plaing. Control lateral movement of NAPL by pressure injecting hydrautic coments, clays, bentonite, and silicates into the formation through tightly spaced borrings using jetting tools. Het Water Injection Control lateral movement of NAPL by pressure injecting hydrautic coments, clays, bentonite, and silicates into the formation through tightly spaced borrings using jetting tools. Het Water Injection Control lateral movement of NAPL by pressure injecting hydrautic coments, clays, bentonite, and silicates into the formation through tightly spaced borrings using jetting tools. Het Water Injection Conductive Heating Thermal Conductive Thermal Conductive Heating Thermal Con | | | addressing soil disturbance and/or | used in conjunction with other technology approaches. |
| Section Sect | | | Slurry Wall | |
| Available carments, clays, bentonite, and silicates into the formation through tightly spaced borings using jetting tools. | <i>In-Situ</i> Containment | Vertical Barriers | Sheet Pile Wall | |
| Low-Temperature Thermal Treatment Thermal Conductive Heating | | | Grout Curtain | hydraulic cements, clays, bentonite, and silicates into the |
| Temperature Thermal Treatment Thermal Treatment Thermal Conductive Heating | | Low- | Hot Water Injection | boiling point of water, increasing the mobility and solubility of |
| Thermal Conductive Heating Steam Injection | | Section Process Options Process Options | pumping from wells, and contaminants are treated. Heating | |
| Mid- Temperature Thormal Treatment Electrical Resistance Heating Electrical Resistance Heating Electrical Resistance Heating Electrical Resistance Heating Electrical Resistance Thormal Conductive Heating Electrical Resistance Electrical Resistance Electrical Resistance Electrical Resistance Heating Electrical Resistance Electrical Resistance Heating Electrical Resistance Electrical Resistance Heating Electrical Resistance | | Technology | resistance when voltage is applied between subsurface | |
| Electrical Resistance Heating Treatment Thermal Treatment | | Temperature Thermal Treatment | Steam Injection | point of water, volatilizing or destroying (by pyrolysis) volatile |
| Thermal Conductive Heating Thermal Conductive Heating | | | | soil vapor extraction, contaminated liquids are removed by pumping from wells, and contaminants are treated. Heating can be performed by injecting steam in vertical wells, thermal |
| Heating High- Temperature Thermal Treatment Thermal Conductive Heating Solidification/ Stabilization Chemical Oxidation Chemical Oxidation Chemical Oxidation Chemical Oxidation Chemical Oxidation Chemical Oxidation Treatment Chemical Oxidation Chemical Oxidation Chemical Oxidation Treatment NAPL Pumping Pumping of NAPL from wells and trenches Surfactant Enhanced Recovery Excavation Excavation Excavation Co-Burning Co-Burning Co-Burning Co-Burning Coff-Site Management Treatment Treatment Treatment Thermal Thermal Disposal Treatment Thermal T | | | | resistance when voltage is applied between subsurface |
| Temperature Thermal Treatment Thermal Treatment Thermal Treatment Thermal Conductive Heating Thermal Conductive Heating Thermal Conductive Heating Thermal Conductive Heating Solidification/ Stabilization Solidification/ Stabilization include polymers, pozzolasns, and cement. Amendments can be mixed with soil in situ using large-diameter augers, soil mixers, or similar equipment. Chemical oxidation involves the injection of chemical oxidants into the subsurface to react with and destroy organic contaminants. Common oxidants include hydrogen peroxide, potassium permanganate, ozone, and sodium persulfate. Pumping of NAPL from wells and trenches Surfactants are injected near NAPL zones in groundwater to mobilize the NAPL, and then the mobilized NAPL is extracted. May be applied with injection-withdrawal technique or with recirculating system. Ex-Situ Treatment Thermal Off-Site Management Off-Site Management Treatment Off-Site NAPL Disposal of recovered Treatment of NAPL via incineration at a hazardous waste | <i>In-Situ</i> Treatment | | | point of water, volatilizing or destroying (by pyrolysis) volatile |
| Stabilization Stabilization Solidification/ Stabilization Solidification/ Stabilization Solidify or immobilize contaminants. Potential amendments include polymers, pozzolans, and cement. Amendments can be mixed with soil in situ using large-diameter augers, soil mixers, or similar equipment. Chemical Treatment Chemical oxidation Chemical oxidation Chemical oxidation Chemical oxidation involves the injection of chemical oxidants into the subsurface to react with and destroy organic contaminants. Common oxidants include hydrogen peroxide, potassium permanganate, ozone, and sodium persulfate. Pumping of NAPL from wells and trenches Pumping to remove NAPL that accumulates in a well or trench. Surfactants are injected near NAPL zones in groundwater to mobilize the NAPL, and then the mobilized NAPL is extracted. May be applied with injection-withdrawal technique or with recirculating system. Excavation Excavation Co-Burning Combustion of coal tar- or tar-contaminated soil with coal in utility boilers and cement kilns. When soil or sediment containing NAPL is heated to temperatures above 1,400°F, contaminants are directly oxidized. Recycling of recovered NAPL Disposal of recovered Treatment of NAPL via incineration at a hazardous waste | | Temperature Thermal | | are removed by pumping from wells, and contaminants are treated. Heating can be performed by thermal conduction from vertical heated wells, or by electrical resistance when voltage is |
| Chemical Treatment Chemical oxidation Into the subsurface to react with and destroy organic contaminants. Common oxidants include hydrogen peroxide, potassium permanganate, ozone, and sodium persulfate. NAPL Pumping | | Stabilization | | solidify or immobilize contaminants. Potential amendments include polymers, pozzolans, and cement. Amendments can be mixed with soil <i>in situ</i> using large-diameter augers, soil |
| Removal Surfactant Enhanced Recovery Excavation Ex-Situ Treatment Disposal NAPL Pumping of wells and trenches Wells and trenches Wells and trenches Pumping of mobilized NAPL Pumping of mobilized NAPL Pumping of mobilized NAPL Pumping of mobilized May be applied with injection-withdrawal technique or with recirculating system. Excavation Co-Burning Co-Burning NAPL is removed by excavating soil containing NAPL. Combustion of coal tar- or tar-contaminated soil with coal in utility boilers and cement kilns. When soil or sediment containing NAPL is heated to temperatures above 1,400°F, contaminants are directly oxidized. Recycling of recovered NAPL Disposal of recovered Treatment of NAPL via incineration at a hazardous waste | | | Chemical oxidation | into the subsurface to react with and destroy organic contaminants. Common oxidants include hydrogen peroxide, |
| Surfactant Enhanced Recovery Pumping of mobilized NAPL mobilize the NAPL, and then the mobilized NAPL is extracted. May be applied with injection-withdrawal technique or with recirculating system. | | NAPL Pumping | | Pumping to remove NAPL that accumulates in a well or trench. |
| Ex-Situ Treatment Thermal Thermal Thermal Disposal Co-Burning Co-Burning Co-Burning Co-Burning Combustion of coal tar- or tar-contaminated soil with coal in utility boilers and cement kilns. When soil or sediment containing NAPL is heated to temperatures above 1,400°F, contaminants are directly oxidized. Recycling of recovered NAPL Disposal of recovered Treatment of NAPL via incineration at a hazardous waste | Removal | Enhanced | | mobilize the NAPL, and then the mobilized NAPL is extracted. May be applied with injection-withdrawal technique or with |
| Ex-Situ Treatment Thermal Thermal Thermal Thermal Thermal Thermal Thermal Thermal Incineration Thermal Thermal Incineration Thermal Thermal Incineration Thermal Thermal Incineration Thermal | | Excavation | | |
| Disposal Off-Site Management Recycling of recovered NAPL Reuse of recovered product. Disposal of recovered Treatment of NAPL via incineration at a hazardous waste | Ex-Situ Treatment | Thermal | | utility boilers and cement kilns. When soil or sediment containing NAPL is heated to temperatures above 1,400°F, contaminants are directly |
| Disposal of recovered Treatment of NAPL via incineration at a hazardous waste | Diamond | Off-Site | | |
| NAPL via incineration treatment facility. | ⊔isposal | | Disposal of recovered | |

Notes:

¹Institutional controls are not considered stand-alone remedial technologies but may be applied in conjunction with other cleanup technologies.

BTEX = benzene, tolouene, ethylbenzene, and xylenes

cPAHs = carcinogenic polycyclic aromatic hydrocarbons

NAPL = non-aqueous phase liquid O&M = operation and maintenance

PAHs = polycyclic aromatic hydrocarbons

TPH = total petroleum hydrocarbons

| Soil General Response Actions | Remedial Technology | Process Options | Description |
|--|---------------------------------------|---|--|
| | Access Restrictions | Fences and warning signs to control Site access | Signs, fences, or other measures to prevent access to the property. |
| Institutional Controls ¹ | Use Restrictions | Use restrictions and monitoring to prevent disturbance of engineered controls | Covenant placed on the property that limits or prohibits activities that may interfere with a cleanup action or result in exposure to hazardous substances. |
| | | Deed restrictions addressing soil disturbance | |
| | | Permeable soil cover | Placing clean soil on the surface provides a barrier that prevents exposure to underlying soil but allows storm water to infiltrate. |
| <i>In-Situ</i> Containment | Capping | Low-permeability cap | Low-permeability caps may be constructed of low-permeability soil such as clay or an engineered material such as asphalt or concrete. This cap would not only prevent exposure to underlying soils, but would also minimize stormwater infiltration through potentially contaminated materials, thereby reducing mobility of contaminants located in the unsaturated soil zone. Engineered materials could also be used in areas requiring a durable surface, such as high-traffic areas. |
| | | Impervious cap | Impervious caps may be constructed of low-permeability soil such as clay or an engineered material such as asphalt or concrete, overlain by an additional impermeable layer. This cap would not only prevent exposure to underlying soils, but would also prevent stormwater from infiltrating through potentially contaminated soils beneath the cap, thereby reducing mobility of contaminants located in the unsaturated soil zone. Often combined with barrier wall technology to fully encapsulate soils. |
| | Physical Removal and Treatment | Passive venting of soil vapors | Passive soil venting is a less aggressive version of soil vapor extraction that is usually applied to prevent contaminated soil vapors from migrating into buildings or crawl spaces. In passive venting, soil vapors beneath a building foundation are vented to the atmosphere either through atmospheric pressure changes or by applying a low vacuum with a ventilation fan. Vented vapors can be passed through activated carbon for treatment if necessary. |
| | | Soil vapor extraction | Soil vapor extraction applies a vacuum to subsurface soil to volatilize contamination and extract soil vapor. Vapor stream is treated above ground to remove contamination before discharge. |
| | Low-Temperature Thermal Treatment | Hot Water Injection | The subsurface is heated to temperatures less than the boiling point of water, increasing the mobility and solubility of NAPL and |
| | | Electrical Resistance Heating | NAPL constituents. Contaminated liquids are removed by pumping from wells, and contaminants are treated. Heating can be performed by injecting steam in vertical wells, thermal conduction |
| 1.07 | | Thermal Conductive Heating | from vertical heated wells, or by electrical resistance when voltage is applied between subsurface electrodes. |
| <i>In-Situ</i> Treatment | | Steam Injection | The subsurface is heated to temperatures near the boiling point of water, volatilizing or destroying (by pyrolysis) volatile organic |
| | Mid-Temperature Thermal Treatment | Electrical Resistance Heating | compounds. Contaminated vapors are collected using soil vapor extraction, contaminated liquids are removed by pumping from wells, and contaminants are treated. Heating can be performed by |
| | | Thermal Conductive Heating | injecting steam in vertical wells, thermal conduction from vertical heated wells, or by electrical resistance when voltage is applied between subsurface electrodes. |
| | High-Temperature Thermal Treatment | Thermal Conductive Heating | The subsurface is heated to temperatures above the boiling point of water, volatilizing or destroying (by pyrolysis) volatile and semi-volatile organic compounds. Contaminated vapors are collected using soil vapor extraction, contaminated liquids are removed by pumping from wells, and contaminants are treated. Heating can be performed by thermal conduction from vertical heated wells, or by electrical resistance when voltage is applied between subsurface electrodes. |
| | | Vitrification | Soil is heated via electrical current to temperatures greater than 2,400°F, destroying contaminants and fusing soil into a glassy matrix. |

Table 7-2 - Remedial Technologies for Soil

Bremerton Gas Works Superfund Site Bremerton, Washington

| Soil General Response Actions | Remedial Technology | Process Options | Description |
|-------------------------------------|-------------------------------|--|---|
| | Stabilization | Solidification/ Stabilization | Soil or sediment is stabilized by adding amendments to solidify or immobilize contaminants. Potential amendments include polymers, pozzolans, and cement. Amendments can be mixed with soil <i>in situ</i> using large-diameter augers, soil mixers, or similar equipment. |
| <i>In Situ</i> Treatment | Chemical Treatment | Chemical oxidation | Chemical oxidation involves the injection of chemical oxidants into the subsurface to react with and destroy organic contaminants. Common oxidants include hydrogen peroxide, potassium permanganate, ozone, and sodium persulfate, which have been shown to destroy a wide range of contaminants in soil. |
| | | Bioventing | Bioventing supplies oxygen to unsaturated soil to increase aerobic biodegradation rates and may be designed to increase the air exchange rate through the soil. |
| | Bioremediation | Amendment Injection | Biodegradation of contaminants by indigenous soil microbes can be enhanced by amending soil with nutrients, moisture, and oxygen (typically provided by injecting air or solutions into wells or trenches). |
| Removal | Excavation | Excavation | Excavators, backhoes, and other conventional earth moving equipment are the most common equipment used to remove contaminated soil from upland areas. |
| | Physical | Solidification/ Stabilization | Amendments are added to excavated soil or sediment to immobilize and/or bind contaminants within the stabilized product. Depending on the proportion of amending agents, the end product may take on the form of a quasi-soil/concrete material that could later be used as bulk fill. |
| | Thermal | Co-Burning | Combustion of Manufactured Gas Plant residues, such as coal tar and tar contaminated soil, with coal in utility boilers and cement kilns. |
| | | Thermal desorption | Low-temperature thermal desorption involves heating soils or sediments to temperatures between 200°F and 600°F until volatile and semivolatile chemicals of concern (COCs) such as benzene and naphthalene evaporate. Exhaust gases produced by the process are typically combusted. |
| | | Incineration | When soil is heated to temperatures above 1,400°F, contaminants are directly oxidized. |
| Ex-Situ Treatment | Chemical/ Physical | Particle washing | In particle washing, soil is put in contact with an aqueous solution to remove contaminants from the soil particles. The suspension is often also used to separate fine particles from coarser particles, allowing beneficial use of the coarser fraction (if sufficiently clean) at the Site. |
| | | Solvent extraction | Solvent extraction is a variant of soil washing in which an organic solvent (rather than an aqueous solution) is put in contact with the soil to remove contaminants. |
| | | Landfarming | Microbial population potentially enhanced with nutrients, moisture, and bioaugmentation to treat contaminated soil on lined beds with tilling and irrigation. |
| | Bioremediation | Biopiles | Microbial population potentially enhanced with nutrients, moisture, aeration, and bioaugmentation to treat contaminated soil in stockpiles. |
| | | Bioreactor | Microbial population potentially enhanced with nutrients, moisture, aeration, and bioaugmentation to treat contaminated soil in enclosed reactor vessels. |
| | | Cold-Mix Asphalt Batching | Encapsulation of contaminant by blending residues, wet aggregate and asphalt emulsion at ambient temperature. |
| Reuse | Asphalt Batching | Hot-Mix Asphalt Batching | Encapsulation of contaminant by blending residues, wet aggregate and asphalt emulsion at high temperature. |
| Disposal | Confined On-Site Disposal | Confined On-site disposal | Excavated soils exceeding applicable cleanup standards could potentially be placed on site in a specially designed upland confined disposal facility (CDF). Depending on the leachability of confined materials, the CDF could potentially include a liner and a liquid collection system to prevent leachate from contaminating groundwater. |
| | Off-Site Landfill Disposal | Subtitle D (Solid Waste) Subtitle C (Hazardous Waste) | Contaminated soils from the Site may be transported to an off-site, permitted disposal facility. This disposal method provides for secure, long-term containment of hazardous and non-hazardous solid wastes. |

Notes:

¹Institutional controls are not considered stand-alone remedial technologies but may be applied in conjunction with other cleanup technologies.

BTEX = benzene, tolouene, ethylbenzene, and xylenes

cPAHs = carcinogenic polycyclic aromatic hydrocarbons

NAPL = non-aqueous phase liquid

O&M = operation and maintenance

PAHs = polycyclic aromatic hydrocarbons

TPH = total petroleum hydrocarbons

Table 7-3 - Remedial Technologies for Groundwater Bremerton Gas Works Superfund Site

| Groundwater General Response Actions | Remedial Technology | Process Options | Description |
|---|----------------------------------|--|--|
| Institutional | Deed Restrictions | Deed restrictions to preclude drinking water use | Covenant placed on property that limits or prohibits activities that may interfere with a cleanup action or result in exposure to hazardous |
| Controls ¹ | 2004 ((004)040110 | Deed restrictions addressing groundwater wells | substances. |
| Monitored Natural Attenuation | Monitored Natural Attenuation | Groundwater Monitoring | Provides monitoring to document the presence and effectiveness of natural processes in removing or containing Site chemicals of concern (COCs). |
| | | Slurry Wall | Control lateral movement of contaminated groundwater by installing impermeable vertical barriers. Vertical barriers can be constructed of a |
| | Vertical Barriers | Sheet Pile Wall | variety of materials and installation techniques, including driving or vibrating steel sheet piling, excavation of a trench and backfilling with a low-permeability material (e.g., bentonite slurry), in situ mixing of |
| | | Grout Curtain | bentonite with native soils, or pressure injecting hydraulic cement and bentonite. |
| <i>In-Situ</i> Containment | Pumping | Pumping from vertical wells or trenches | Migration of contaminants dissolved in groundwater can be controlled by pumping groundwater from vertical wells or trenches, creating a capture zone within which groundwater flows toward the capture point. |
| | Stormwater Controls | Targeted Infiltration | A hydraulic barrier can be created by collecting and infiltrating stormwater and forming a local groundwater "mound." |
| | | Reduced Infiltration | Hydraulic controls can reduce localized infiltration and seepage of stormwater in impacted areas along the shoreline. |
| | Permeable Reactive Barrier | Sorptive/Reactive Wall | A 40-foot-deep trench may be excavated in the uplands and filled with a permeable material that sorbs dissolved-phase contaminants, facilitating further biodegradation and limiting contaminant migration toward marine sediment and surface water and offshore groundwater. A shallow trench could also excavated on the beach near the shoreline, but would be impacted by brackish water and tidally-influenced groundwater gradients. |
| <i>In-Situ</i> Treatment | Chemical Treatment | Chemical Oxidation | Chemical oxidation involves the injection of oxidant solutions into saturated groundwater to react with and destroy organic contaminants. Common oxidants include hydrogen peroxide, potassium permanganate, ozone, and sodium persulfate. |
| | | Amendment Injection | Injecting compounds, such as peroxides, oxygen-releasing compound, or nutrients, that enhance degradation of contaminants. |
| | Bioremediation | Biosparging | Biosparging involves the injection of oxygen, and sometimes nutrients, to groundwater to enhance aerobic bioattenuation of organic compounds. For volatile contaminants, soil vapor extraction or bioventing may be concurrently applied for unsaturated soil. |
| Removal | Groundwater Extraction | Pumping from Vertical Wells or Trenches | Groundwater can be removed from the subsurface by pumping fluids from wells or trenches. |

Table 7-3 - Remedial Technologies for Groundwater

Bremerton Gas Works Superfund Site Bremerton, Washington

| Groundwater General Response Actions | Remedial Technology | Process Options | Description | |
|---|------------------------|-----------------------------------|--|--|
| | | Adsorption | Granular activated carbon (GAC) can be used to remove organic contaminants. Contaminated groundwater is passed through a bed of GAC, and hydrophobic organic compounds in solution adsorb onto the carbon until the carbon becomes depleted or saturated. Depleted GAC may be regenerated or disposed off Site. | |
| <i>Ex-Situ</i> Treatment | Physical/ Chemical | Air Stripping | Contaminated groundwater and air are typically passed counter- currently through a tower, and volatile contaminants (such as benzene and, to a lesser extent, naphthalene) transfer from the water to the air. The contaminant-laden air is usually treated by activated carbon and then discharged to the atmosphere. | |
| | | Advanced Oxidation Processes | Involves adding chemicals that directly oxidize organic contaminants in water. Process options include ozonation, hydrogen peroxide (with or without catalysts such as Fenton's Reagent or ultraviolet light), and permanganate. | |
| | Biological | Biotreatment | Contaminated groundwater is passed through a biological reactor in which a contaminant-degrading microbial culture is maintained, generally by adding nutrients and oxygen and controlling temperature, pH, and other parameters. Process options include bioslurry reactors, fixed-film bioreactors, and constructed wetlands. | |
| | Off-Site Management | Discharge to Sanitary Sewer | Groundwater is discharged to the local sanitary sewer system. Pretreatment of groundwater may not be required if concentrations of chemicals of concern (COCs) meet discharge criteria. Water containing high concentrations of solids (e.g., from construction dewatering) would likely need to be passed through a settling tank or filter to meet discharge requirements. | |
| Disposal | Management | Discharge to Surface Water | Extracted groundwater may also be discharged to surface water, although this discharge option would likely require a National Pollutant Discharge Elimination System (NPDES) permit. Water discharged to surface water would have to meet strict water quality requirements and would likely require treatment before discharge. | |
| | On-Site Management | Re-introduction to Groundwater | Extracted groundwater may also be discharged on site to groundwater via infiltration galleries or injection wells. Contaminated groundwater would likely require treatment before discharge via this method. | |

Notes:

¹Institutional controls are not considered stand-alone remedial technologies but may be applied in conjunction with other cleanup technologies.

BTEX = benzene, tolouene, ethylbenzene, and xylenes cPAHs = carcinogenic polycyclic aromatic hydrocarbons NAPL = non-aqueous phase liquid O&M = operation and maintenance PAHs = polycyclic aromatic hydrocarbons TPH = total petroleum hydrocarbons

Table 7-4 - Remedial Technologies for Sediment Bremerton Gas Works Superfund Site

| Sediment General | Remedial | Due | Danasis di su | |
|--|-------------------------------|---|--|--|
| Response Actions | Technology | Process Options | Description | |
| | | Governmental advisories and public outreach on fish/shellfish consumption | | |
| Institutional Controls ¹ | Use Restrictions | Easements or restrictive covenants to limit activities which may damage the remedy or increase the potential for exposure | Institutional controls are measures undertaken to limit or prohibit activities that may interfere with a cleanup action or result in exposure to hazardous substances. | |
| | | Monitoring and notification of waterway users to restrict specific activities to protect the remedy | | |
| | Monitored Natural Recovery | Monitored Natural Recovery | A passive remedial approach which relies on monitoring of ongoing, natural processes (physical, biological, and/or chemical mechanisms) that act together to reduce the risk (bioavailability and/or toxicity) of the Site COCs. Monitoring is required to evaluate the effectiveness and frequently includes multiple lines of evidence. | |
| Monitored Natural Recovery | Enhanced Natural Recovery | Thin-Layer Sand Placement | Thin-layer placement normally accelerates natural recovery by adding a layer of clean sediment over contaminated sediment. The acceleration can occur through several processes, including increased dilution through bioturbation of clean sediment mixed with underlying contaminants. Thin-layer placement is typically different than the <i>in situ</i> isolation caps, because it is not designed to provide long-term isolation of contaminants from benthic organisms. | |
| <i>In-Situ</i> Containment | Capping (Non- reactive) | Engineered Sand Cap | An engineered sand cap consists of a layer of granular material placed over contaminated sediments to contain and isolate them from the biologically active surface zone. Engineered caps may also include erosion protection or stability layers such as geosynthetics or armoring materials. | |
| | | Post-Dredge Residuals Management Layer | Similar to cap placement methods described above, with the exception that granular material is applied after dredging to manage residual contamination resulting from dredging. In some cases, a reactive media may be included in the residuals/backfill layer. | |
| <i>In-Situ</i> | Physical/ Chemical | Permeable Reactive Cap | A permeable reactive cap includes a reactive material (such as organoclay, coke, coal, or activated carbon) and similar to a sand cap is placed over contaminated sediments to isolate and contain the contaminated sediments. The reactive material also provides treatment by sorping or binding COCs (dissolved and/or NAPL) and further limiting migration into overlying sediment porewater and surface water. | |
| Treatment | | Stabilization | This technology involves adding amendments to in situ sediment that immobilize and/or bind contaminants within the stabilized media. | |
| | Bioremediation | Amendment Injection | Biodegradation of contaminants by indigenous soil microbes can be enhanced by amending soil with nutrients, moisture, and oxygen (typically provided by injecting into wells or trenches). | |
| Removal | Dredging | Hydraulic | Dredging is the removal of sediment in the wet and is primarily accomplished with hydraulic or mechanical equipment. Hydraulic dredging removes and transports sediment with entrained water in a slurry. Mechanical dredging uses mechanical equipment/force to dislodge and excavate sediment in the wet. Dredging effectiveness may be | |
| кеmoval | Uredging | Dredging Mechanical | | limited by resuspension, release of COCs (i.e., dissolved, particles, and sheens) to water and volatilization to air during dredging, and residual COCs remaining after dredging (USACE 2008). These effects may be reduced by use of containment (e.g., sheet pile, silt curtains) and best management practices. |

Table 7-4 - Remedial Technologies for Sediment

Bremerton Gas Works Superfund Site Bremerton, Washington

| Sediment General Response Actions | Remedial Technology | Process Options | Description | |
|--|---|--|--|--|
| | Physical | Physical Separation | The volume of excavated or dredged contaminated materials may be reduced by physically separating the materials into two or more fractions that can be handled separately. | |
| | · | Stabilization | This technology involves adding amendments to excavated sediment that immobilize and/or bind contaminants within the stabilized media. | |
| Ex-Situ Treatment | Thermal | Thermal Desorption | Low-temperature thermal desorption involves heating soils or sediments to temperatures between 200°F and 600°F until volatile and semivolatile COCs such as benzene and naphthalene evaporate. Exhaust gases produced by the process are typically combusted. | |
| | | Incineration | When sediment is heated to temperatures above 1,400°F, contaminants are directly oxidized. | |
| | On-Site Beneficial Use Confined On-Site Disposal | Sand/Aggregate Reclamation | Dredged material with high sand contents that undergo particle separation may be available for use as concrete aggregate or general upland fill. | |
| | | Topsoil Feedstock | Dredged material may be used as non-organic feedstock for topsoil (i.e., material would be blended with organics). | |
| | | Confined On-site Disposal | Removed sediments exceeding applicable cleanup standards could potentially be placed on Site in a specially designed upland CDF. Depending on the leachability of confined materials, the CDF could potentially include a liner and a liquid collection system to prevent leachate from contaminating groundwater. | |
| Disposal | | Near-shore Confined Disposal Facility (CDF) | Removed sediments exceeding applicable cleanup standards could potentially be placed on Site in a specially designed CDF built along the shoreline. Construction would require significant filling and conversion of aquatic lands. | |
| | | Contained Aquatic Disposal (CAD) | Dredged sediments may be consolidated and disposed of in a deep aquatic excavation adjacent to the Site and capped with clean material. | |
| | Off-Site Landfill | Subtitle D (Solid Waste) | Contaminated sediments from the Site may be transported to an off-Site, permitted disposal facility. This disposal method | |
| | Disposal | Subtitle C (Hazardous Waste) | provides for secure, long-term containment of hazardous an non-hazardous solid wastes. | |

Notes:

¹Institutional controls are not considered stand-alone remedial technologies but may be applied in conjunction with other cleanup technologies.

BTEX = benzene, tolouene, ethylbenzene, and xylenes

COCs = chemicals of concern

cPAHs = carcinogenic polycyclic aromatic hydrocarbons

NAPL = non-aqueous phase liquid

O&M = operation and maintenance

PAHs = polycyclic aromatic hydrocarbons

TPH = total petroleum hydrocarbons

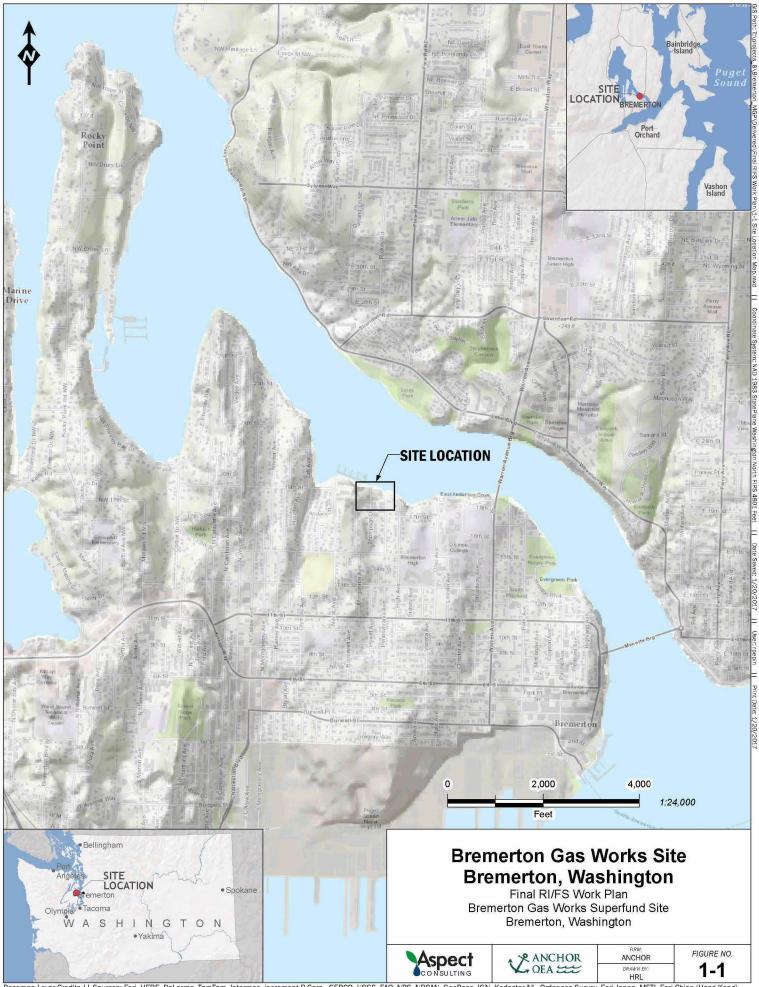
U.S. Army Corps of Engineers (USACE), 2008, Technical Guidelines for Environmental Dredging of Contaminated Sediments, ERDC/EL TR-08-29, September 2008.

Table 8-1 - Estimated Remedial Investigation Data Collection Schedule

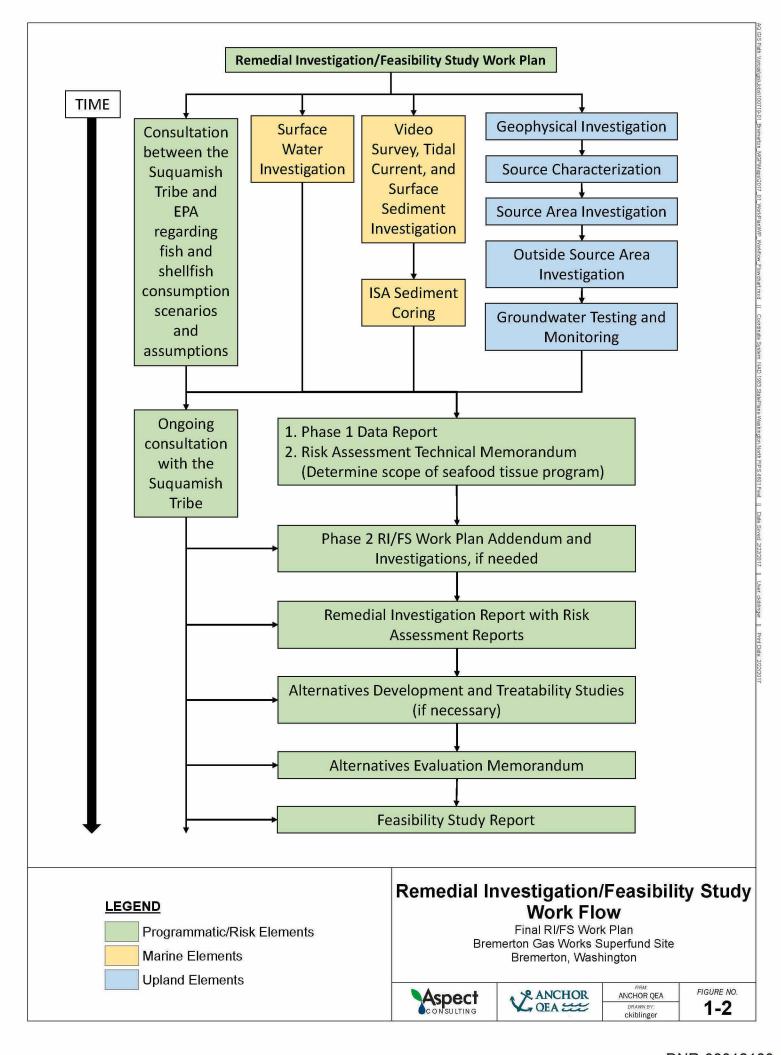
Bremerton Gas Works Superfund Site Bremerton, Washington

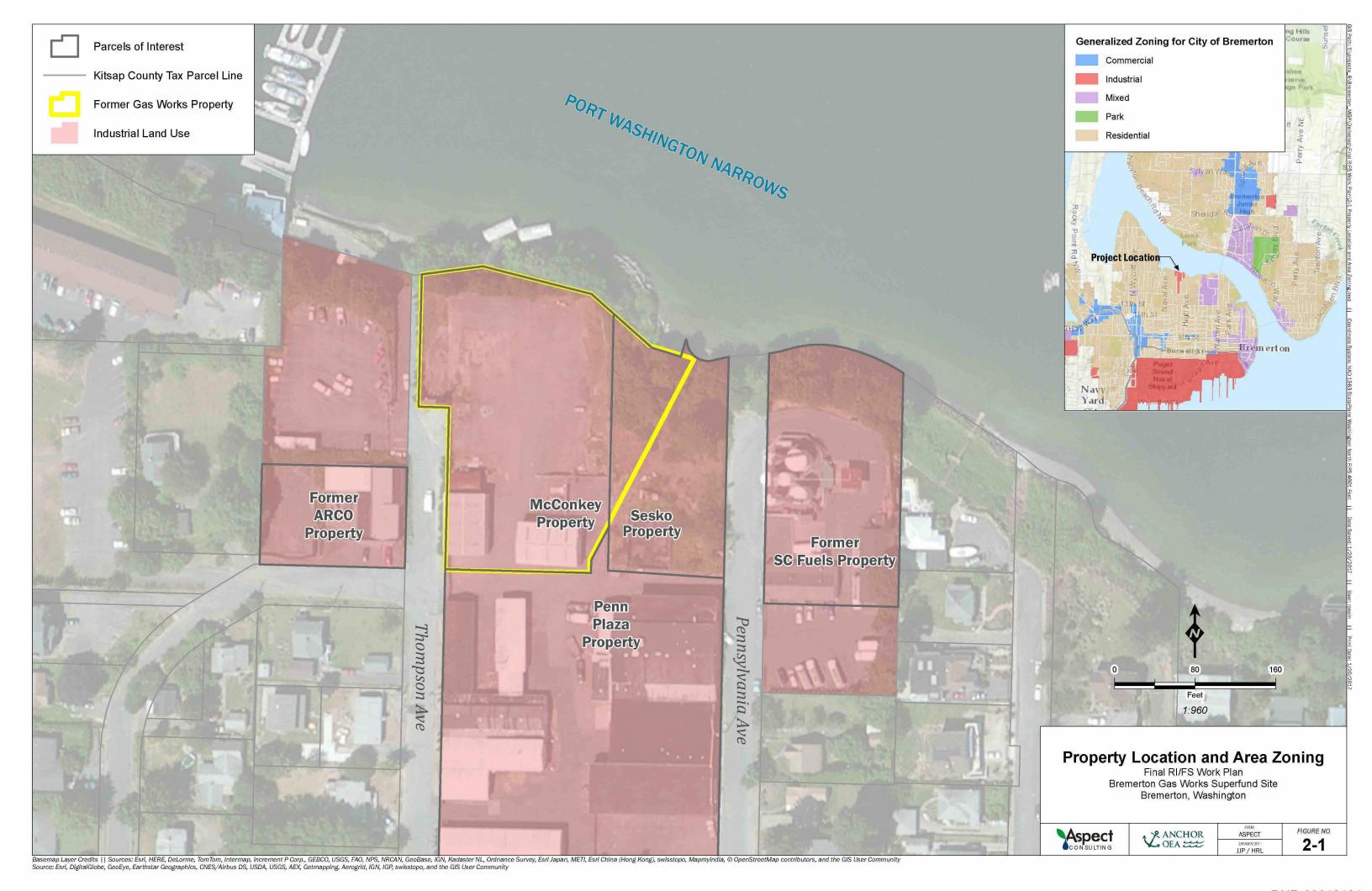
| Task | Estimated Task Duration (Calendar Days) | Time to Completion (Days from Work Plan Approval) | Estimated Completion Date |
|---|---|--|------------------------------|
| Work Plan Approval Date | | | May 1, 2017 |
| Upland Investigation | | | - |
| Contractor coordination and mobilization | 30 | 30 | May 31, 2017 |
| Geophysical Investigation | 15 | 45 | June 15, 2017 |
| Source Characterization | 90 | 135 | September 13, 2017 |
| Source Area Investigation | 120 | 255 | January 11, 2018 |
| Outside Source Area Investigation | 150 | 405 | June 10, 2018 |
| Groundwater Testing and Monitoring | 270 | 675 | March 7, 2019 |
| Marine Area Investigations | | | |
| Towed camera video survey (tidal dependent) | 10 | 365 | May 1, 2018 |
| ADCP Current Survey (tidal dependent) | 2 | 365 | May 1, 2018 |
| Surface Sediment Sampling within ISA (tidal dependent) | 14 | 60 | June 30, 2017 |
| Shellfish Survey (tidal dependent) | 7 | 60 | June 30, 2017 |
| Surface Sediment Sampling within Port Washington Narrows | 7 | 60 | June 30, 2017 |
| Subsurface Sediment Investigation | 10 | 180 | October 28, 2017 |
| Surface Water Sampling | 365 | 365 | May 1, 2018 |
| Phase 1 Data Report (Includes Risk Assessment Tech Memo and W | P Addendum, if applicable | e) | |
| Phase I Data Report | 90 | 765 | June 5, 2019 |

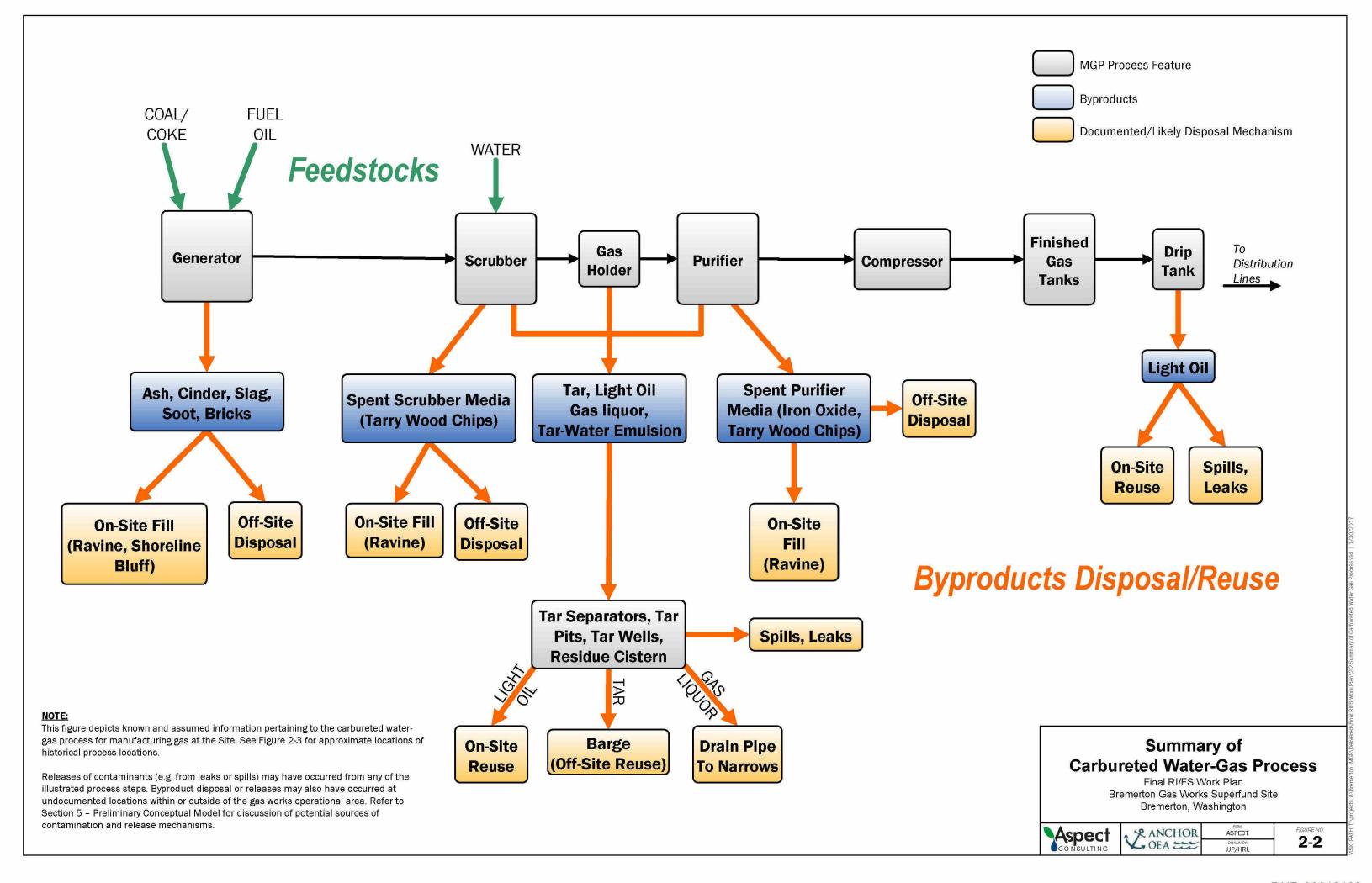
FIGURES

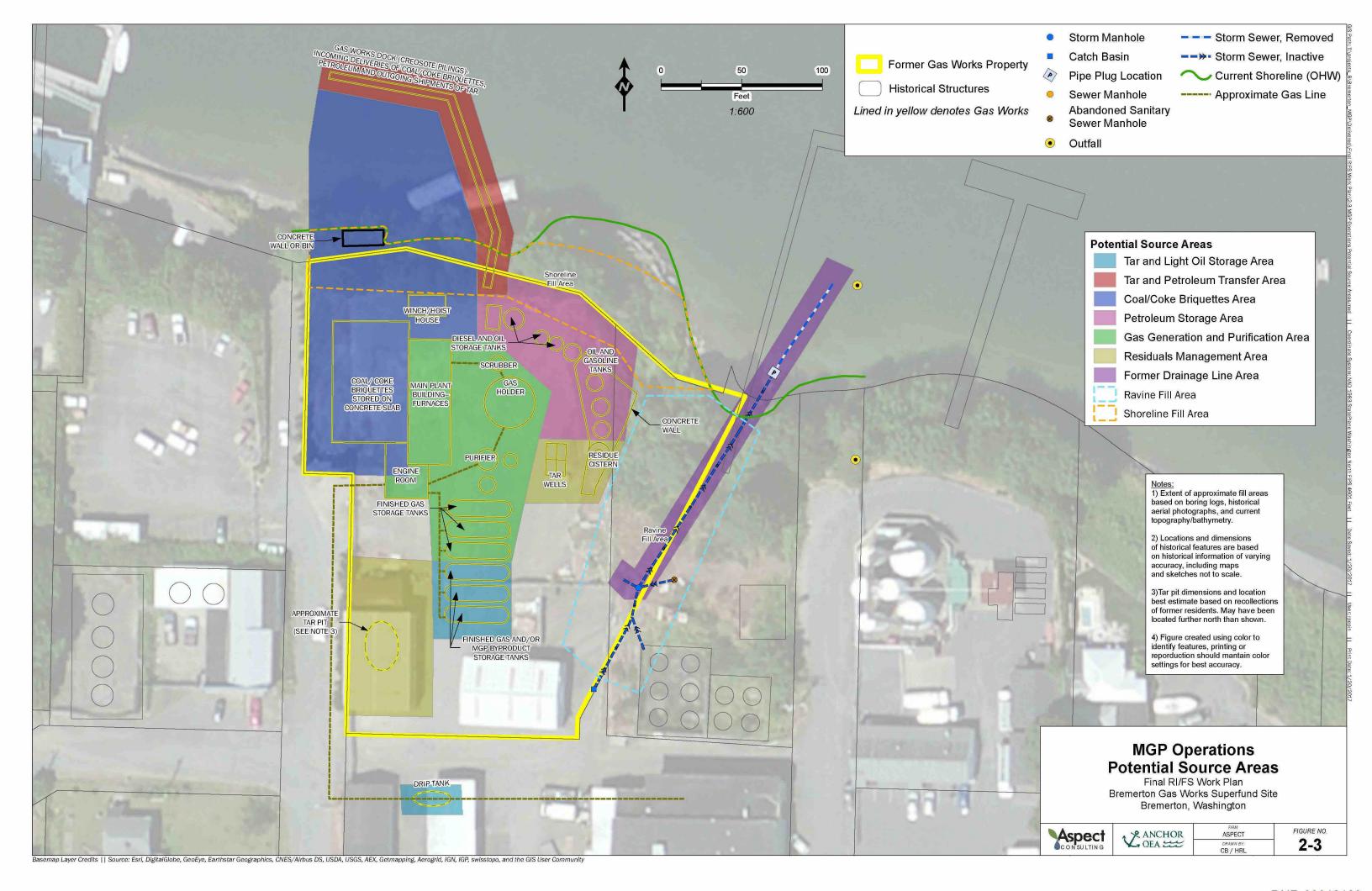


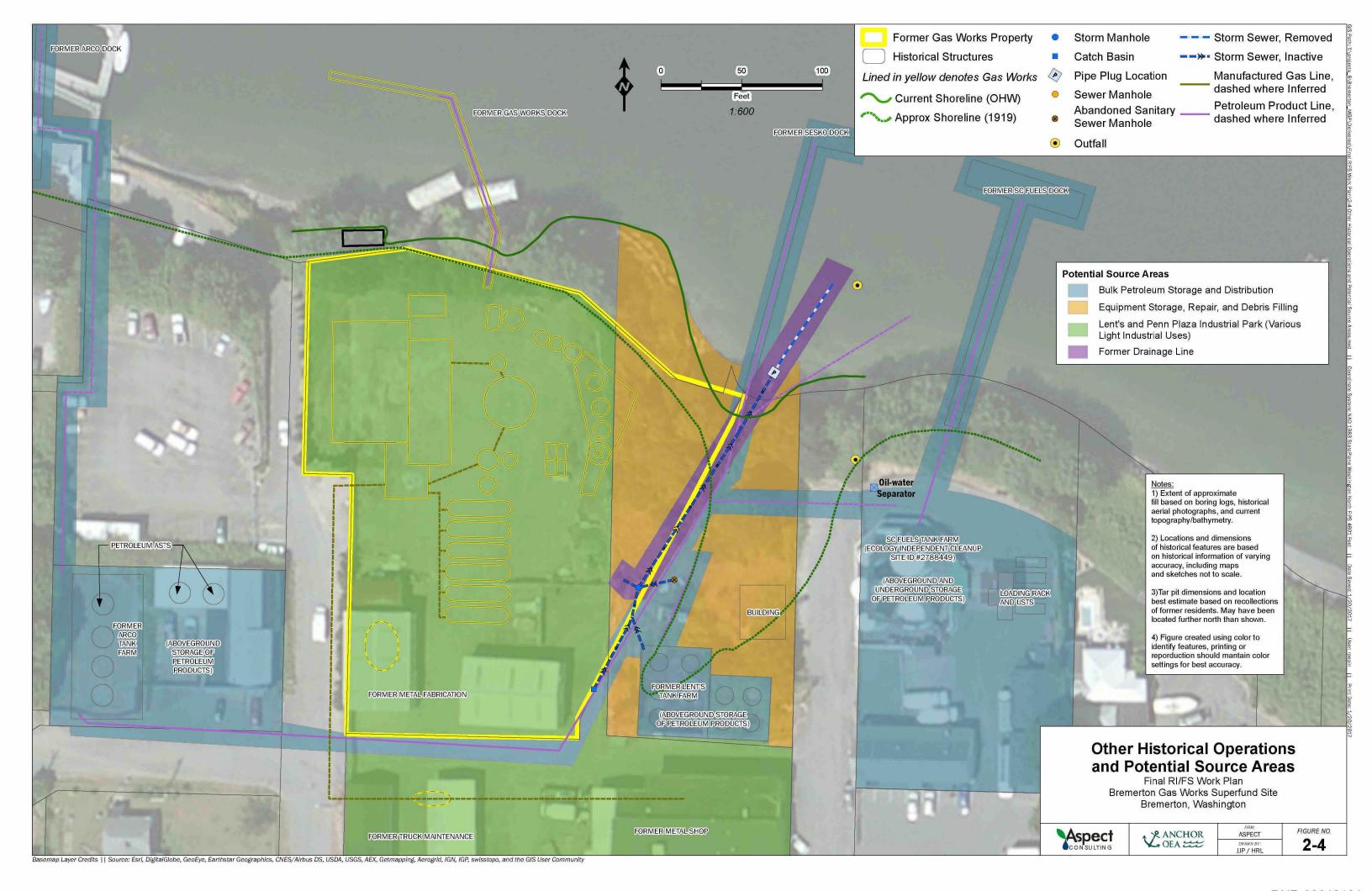
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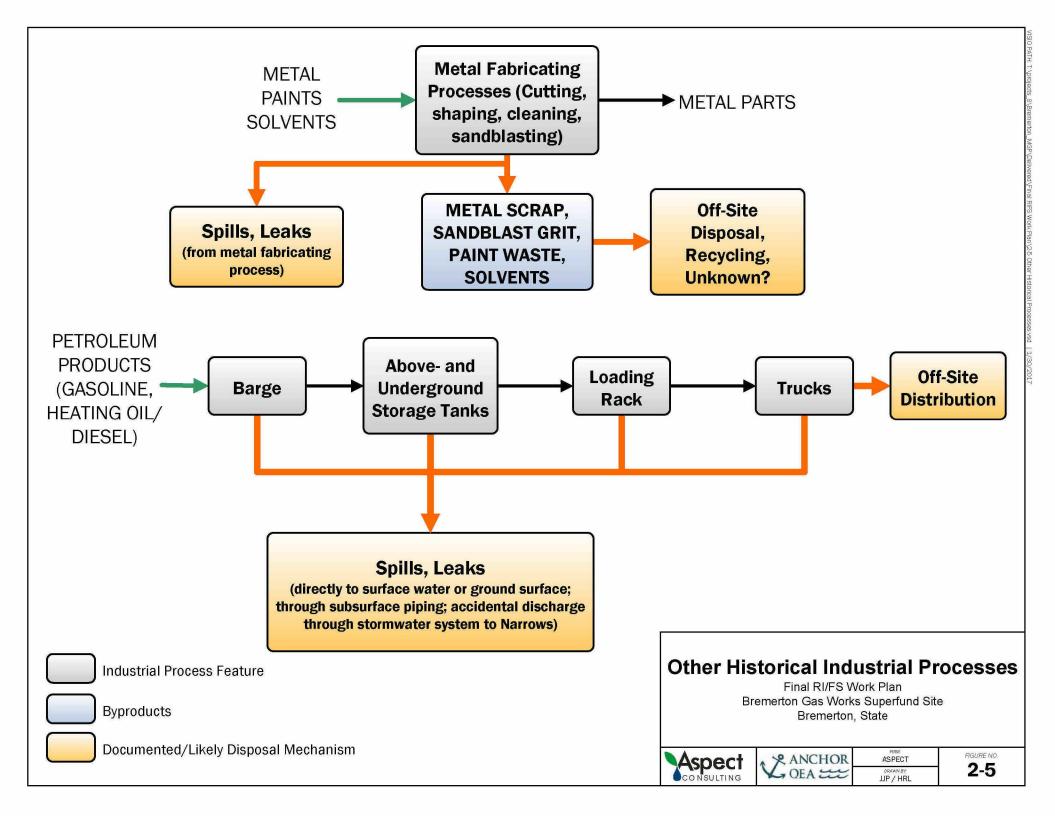


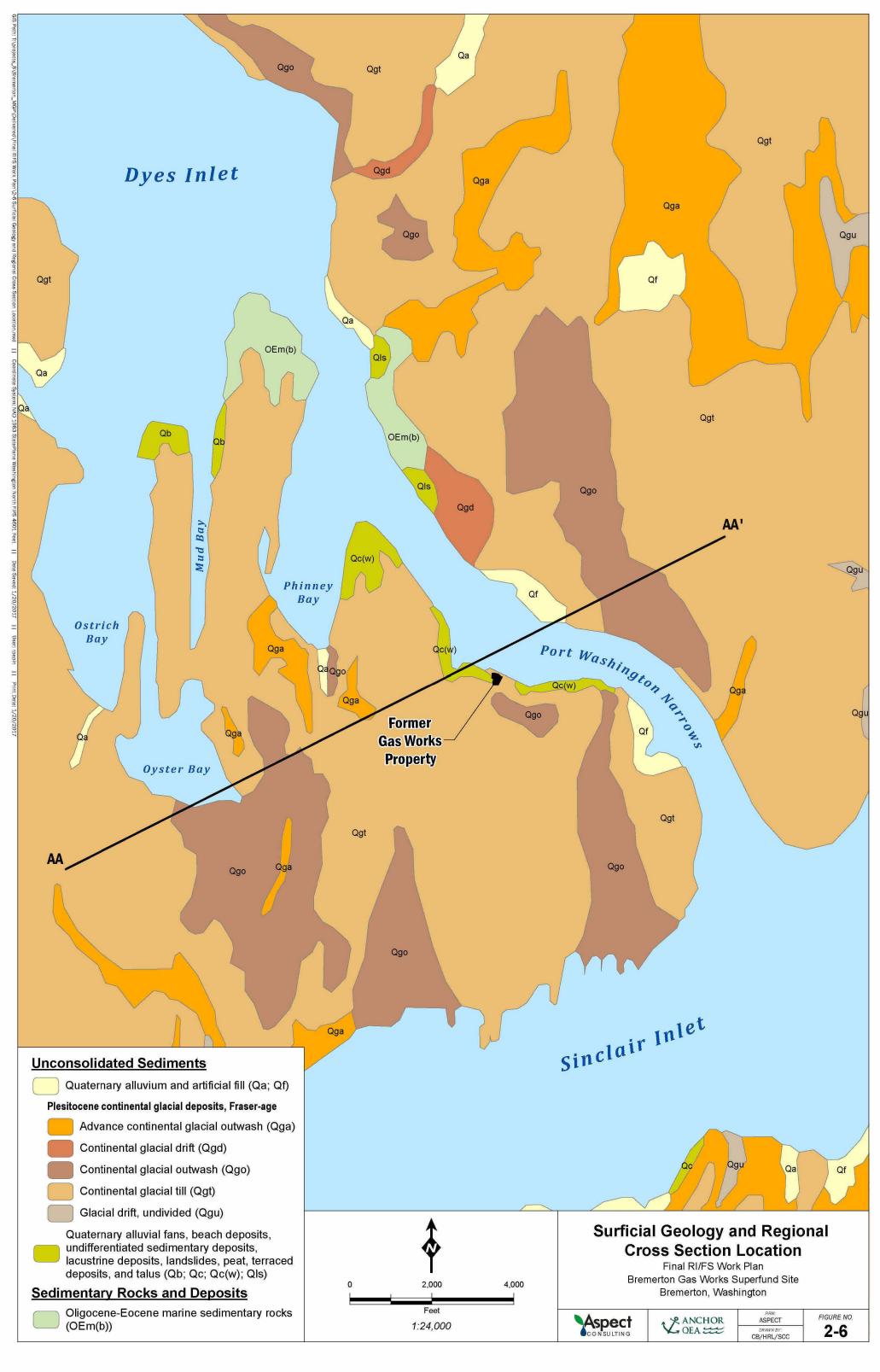


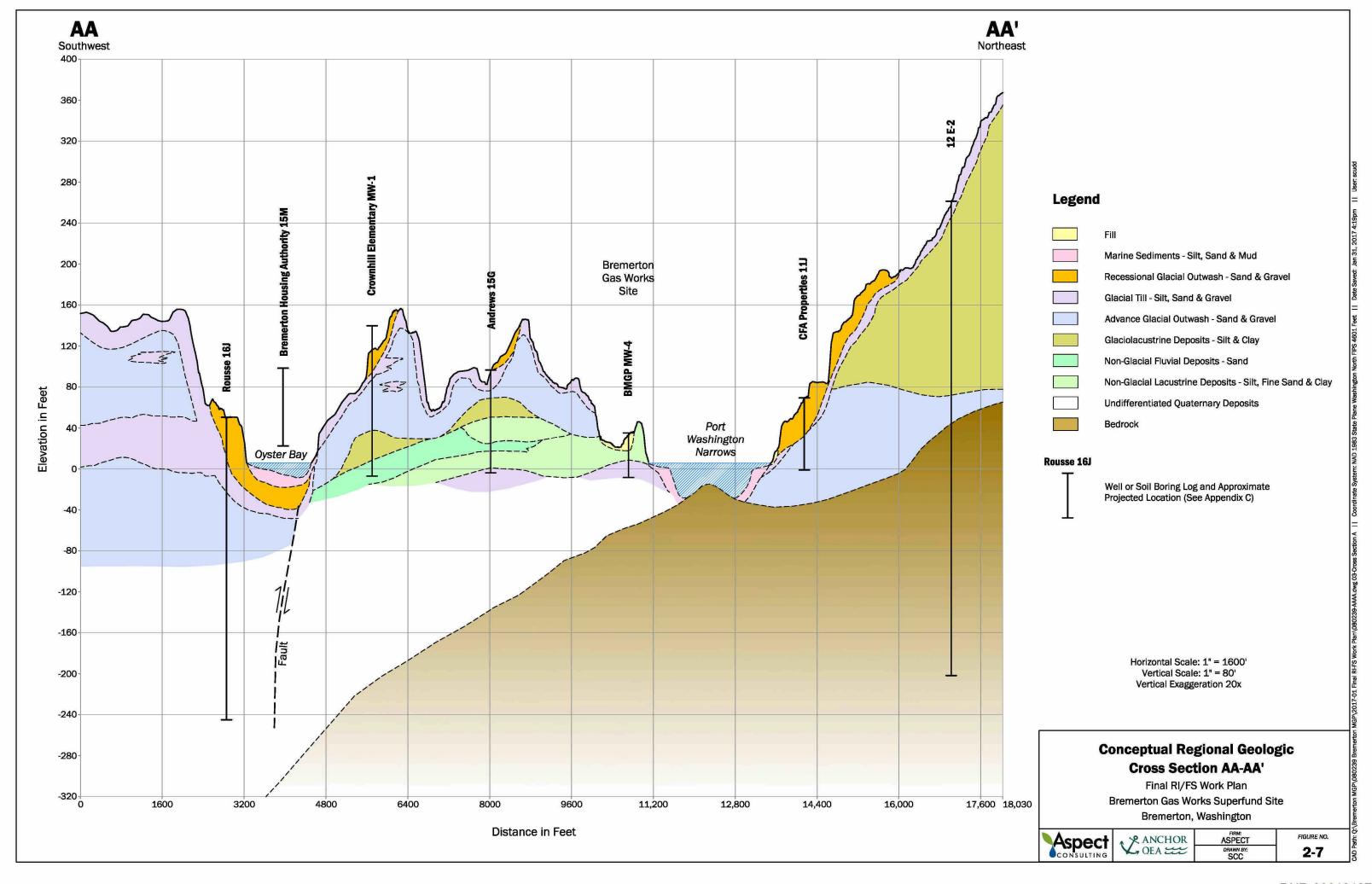


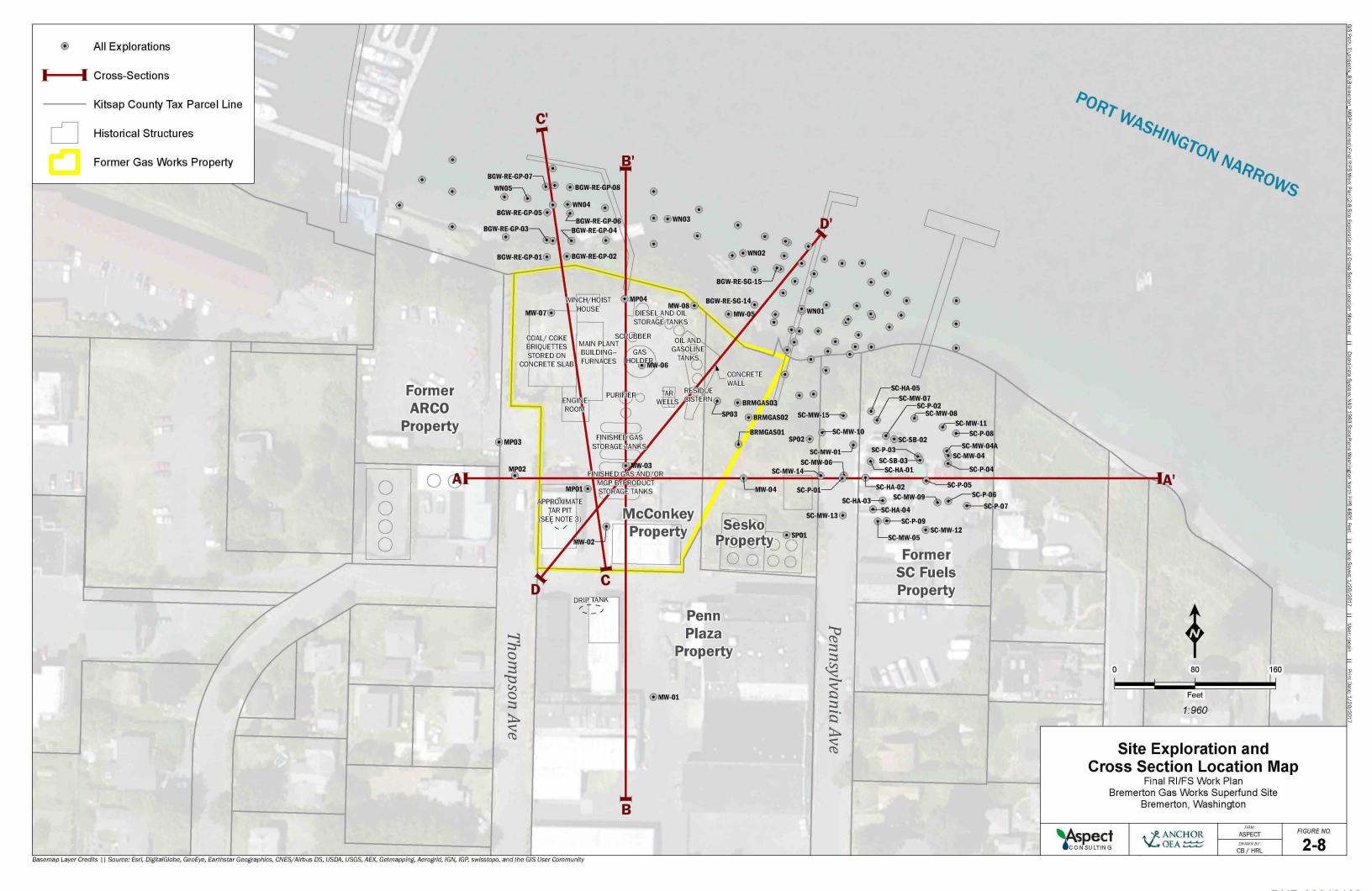


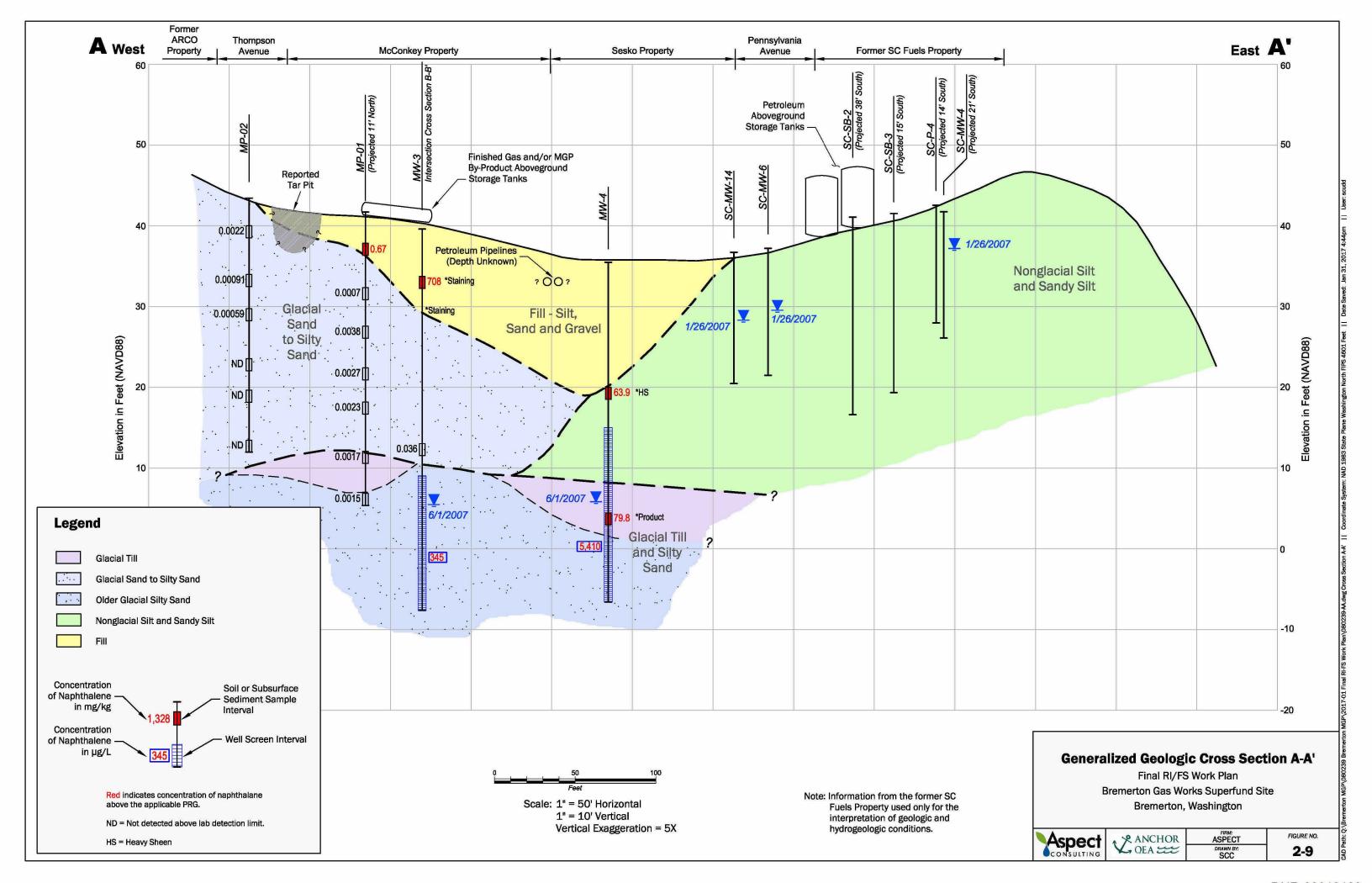


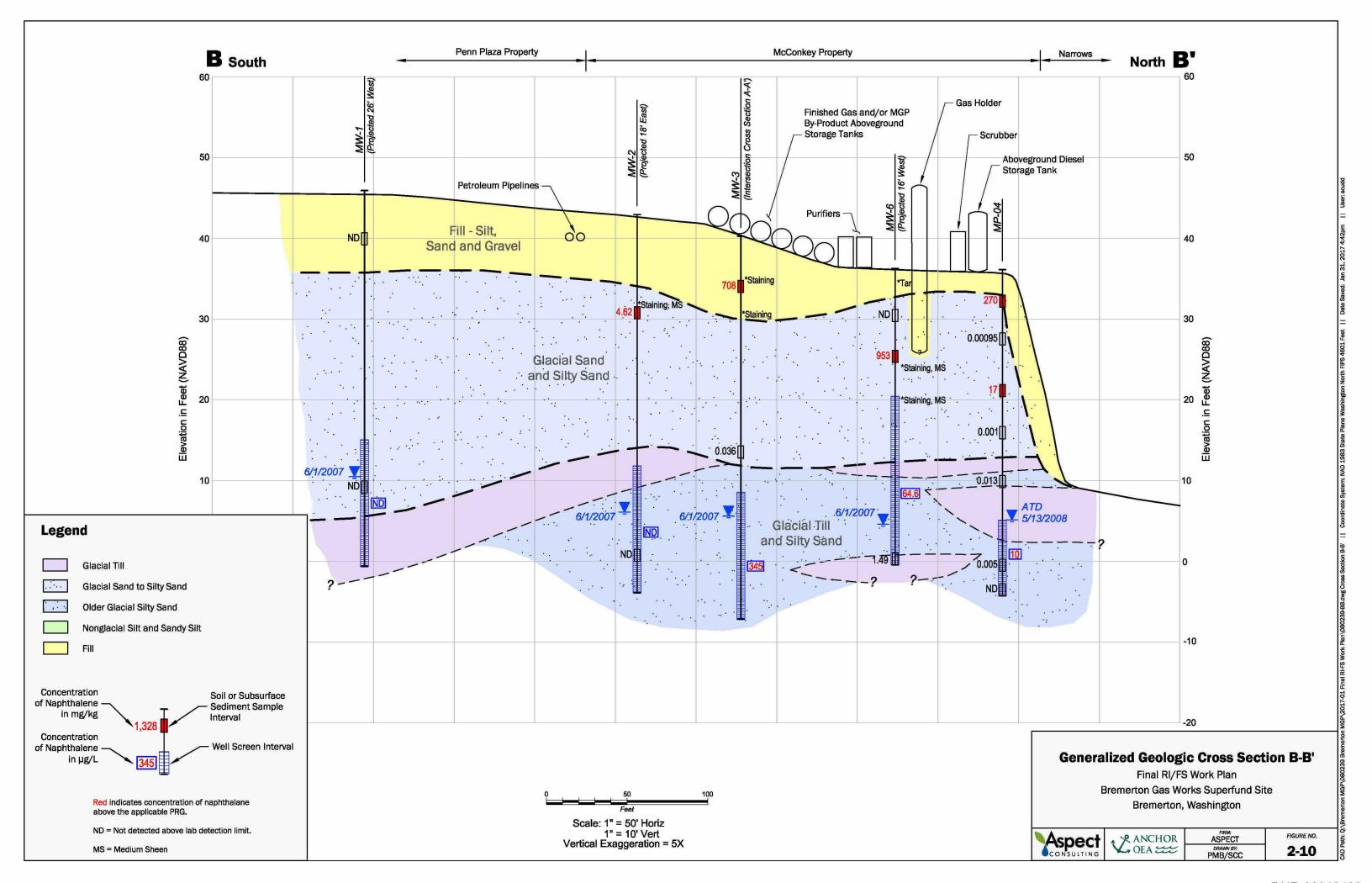


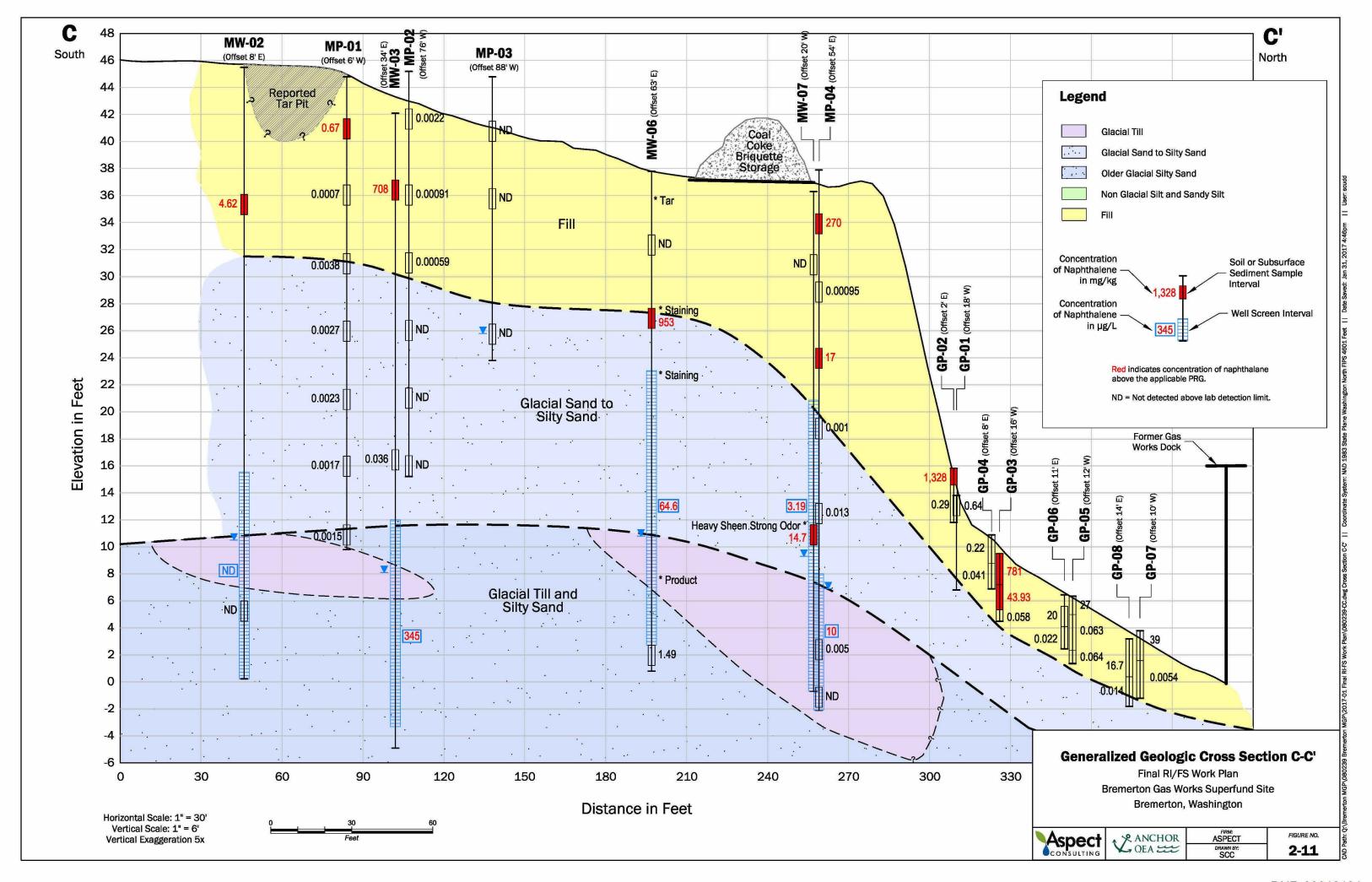


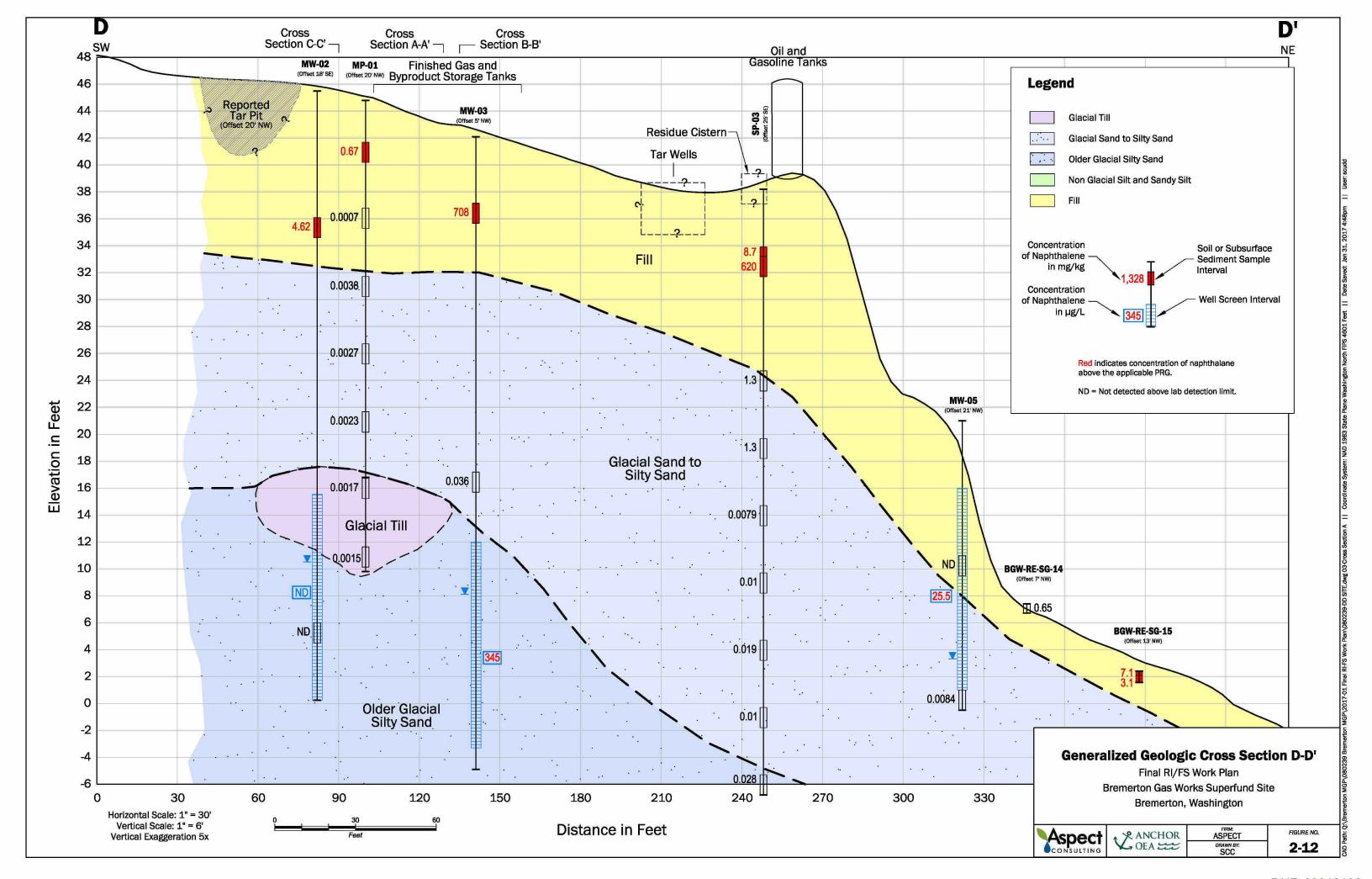


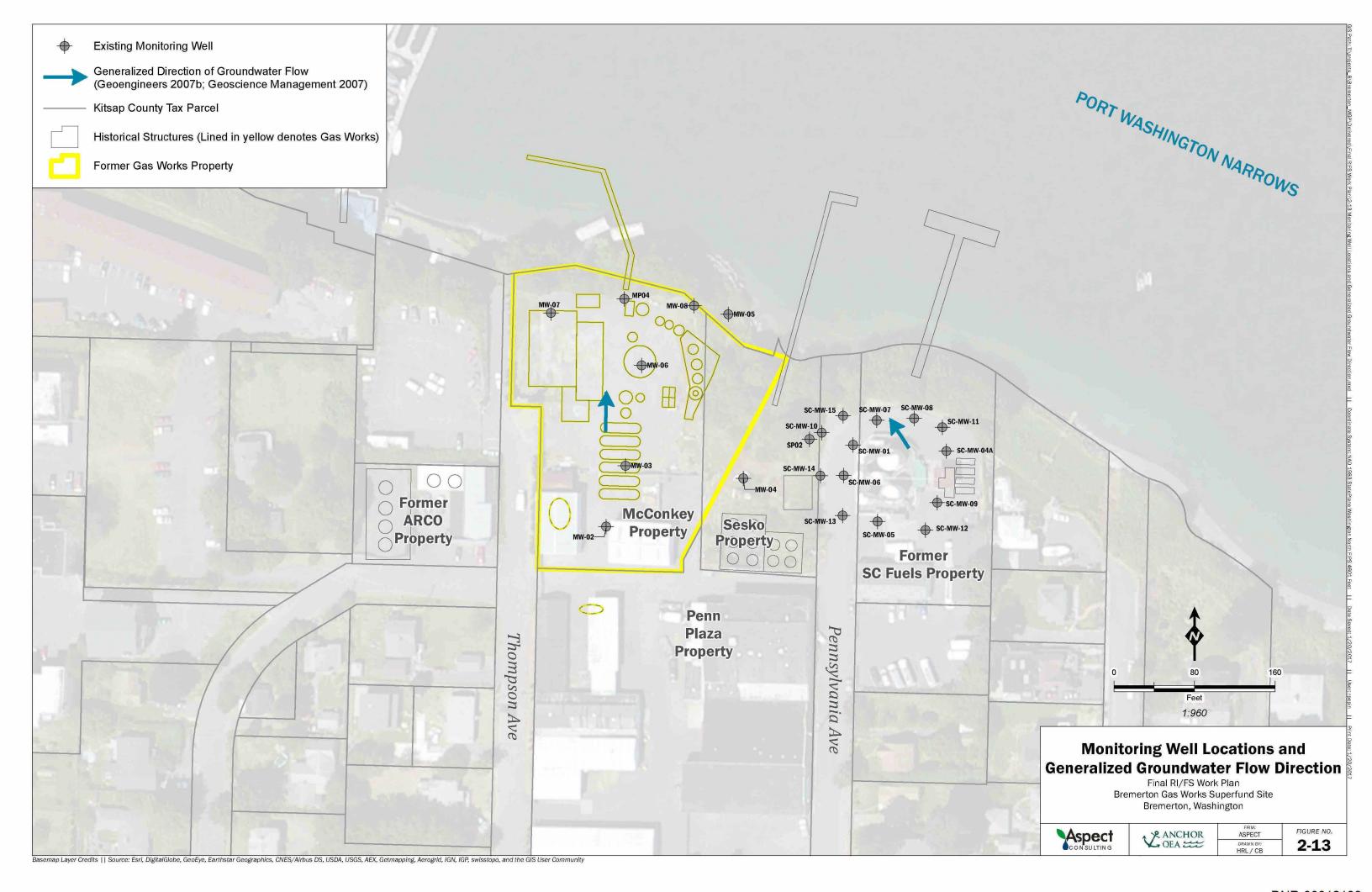


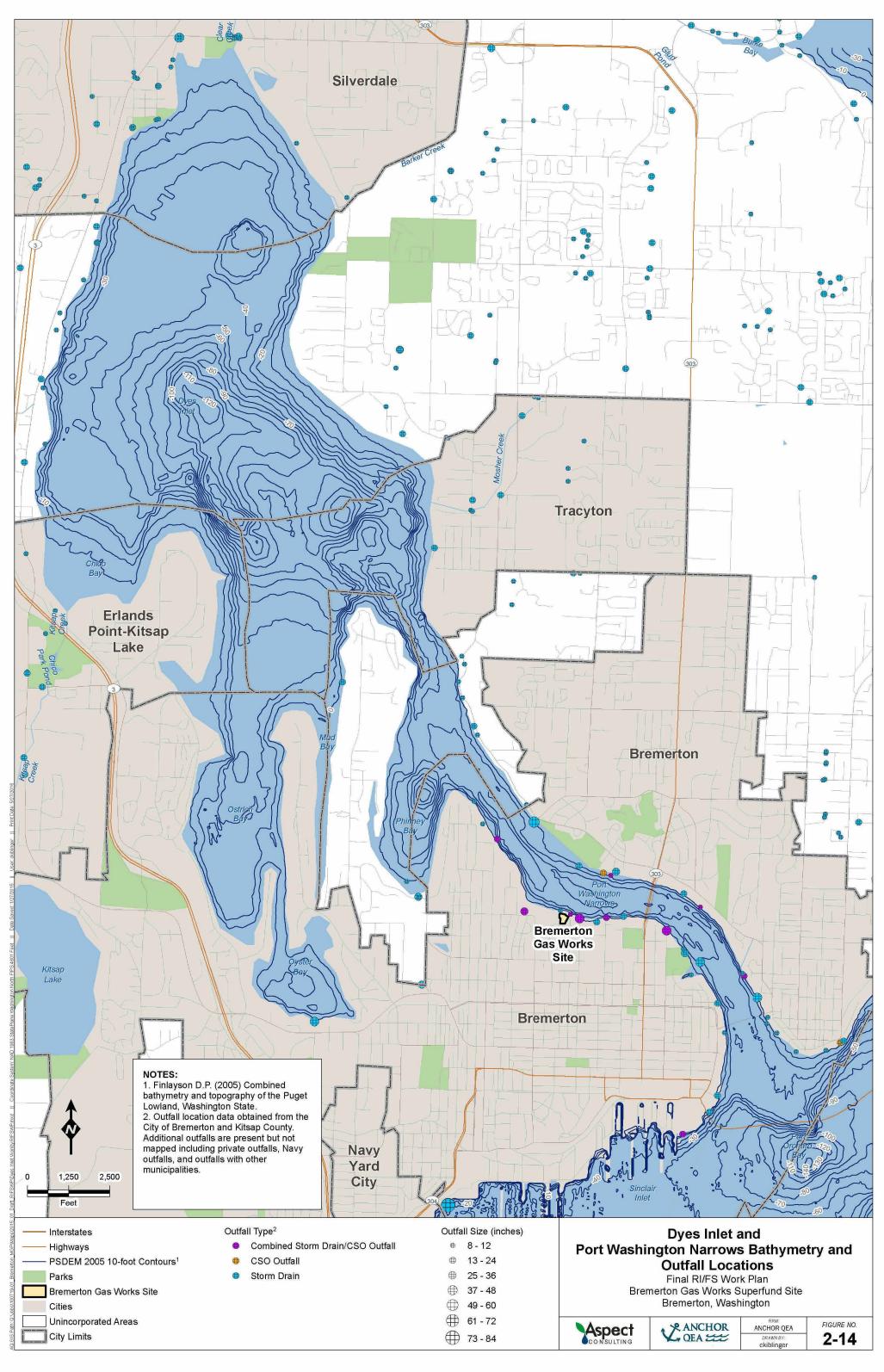


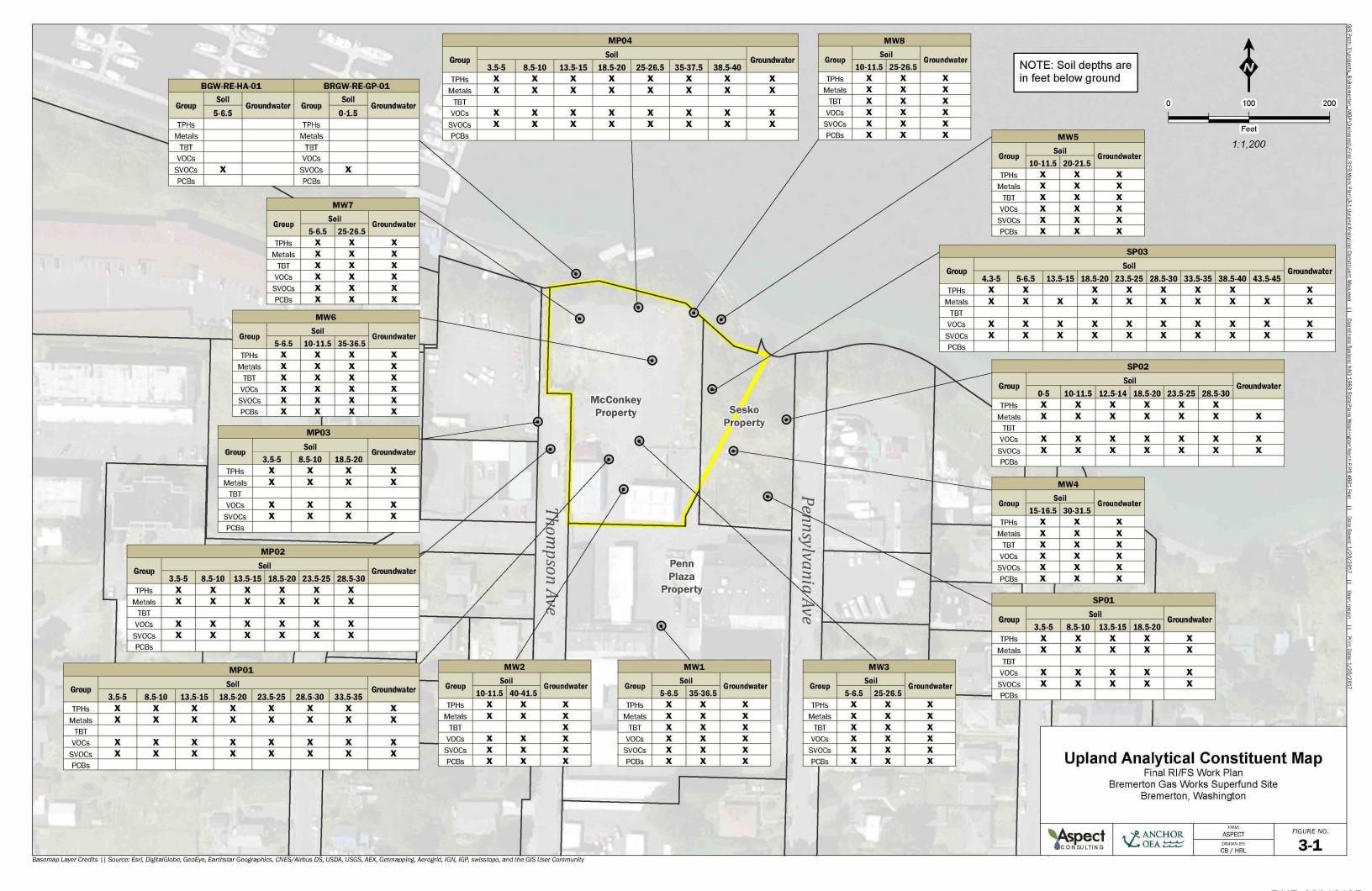


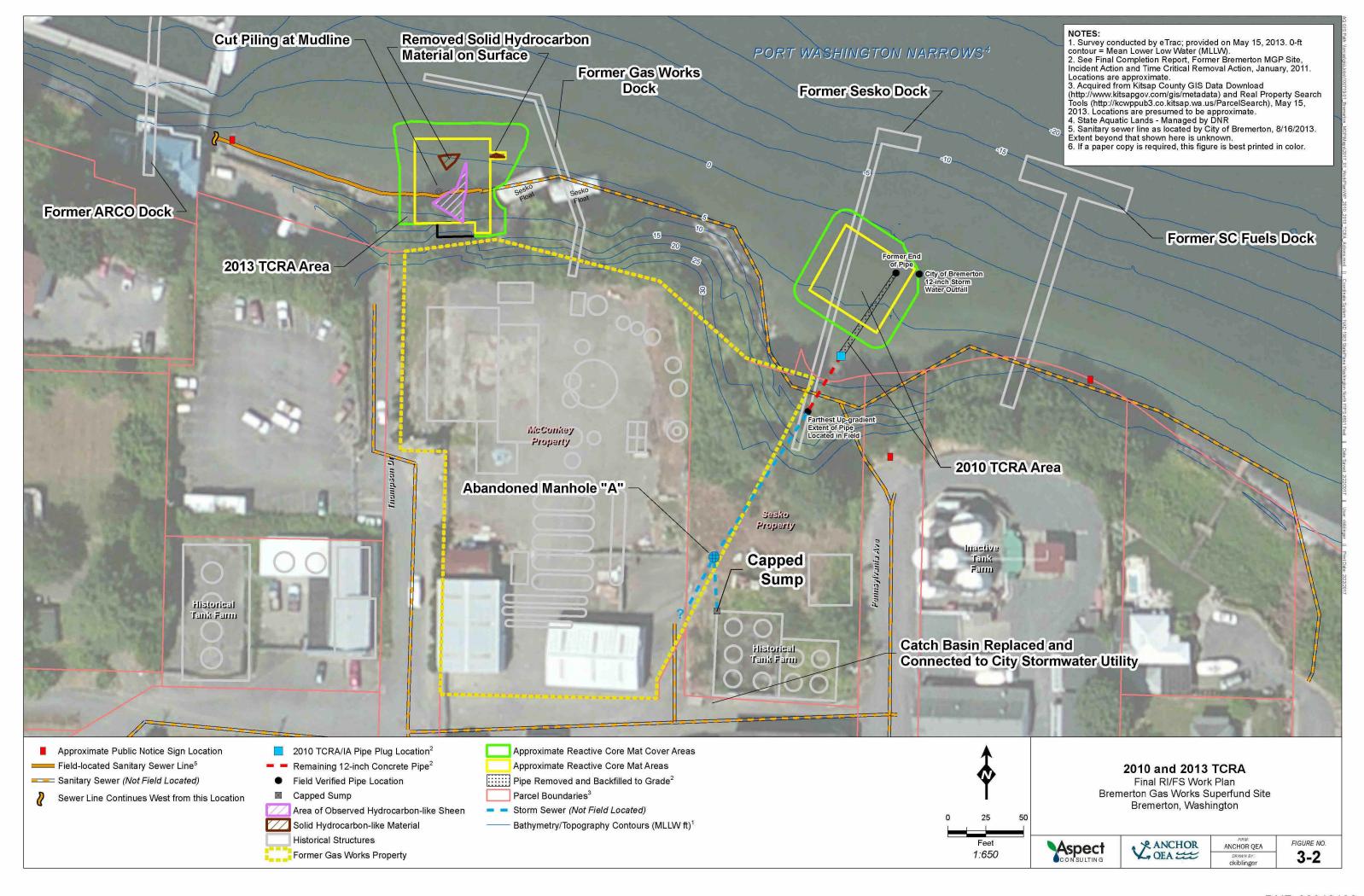


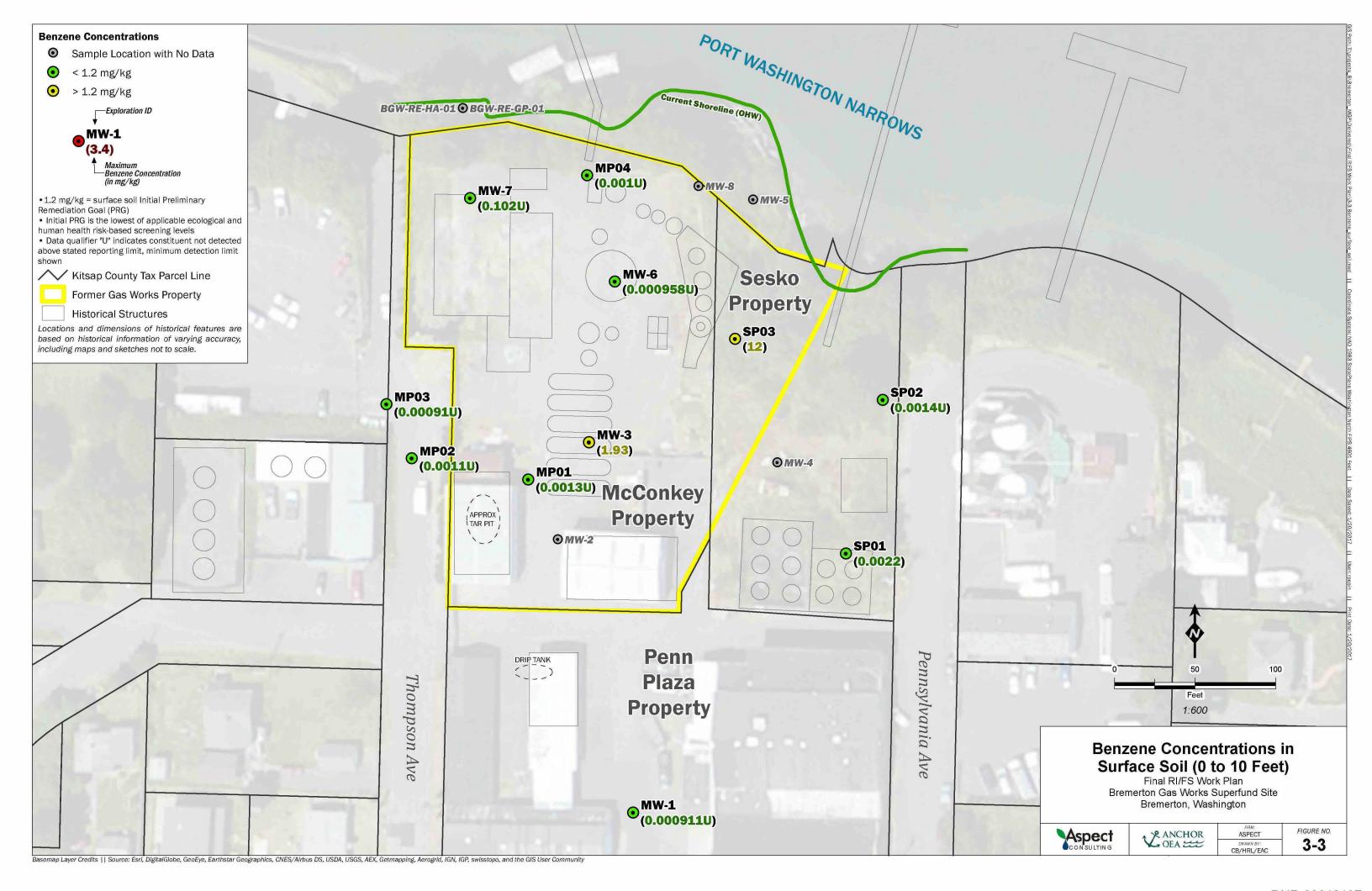


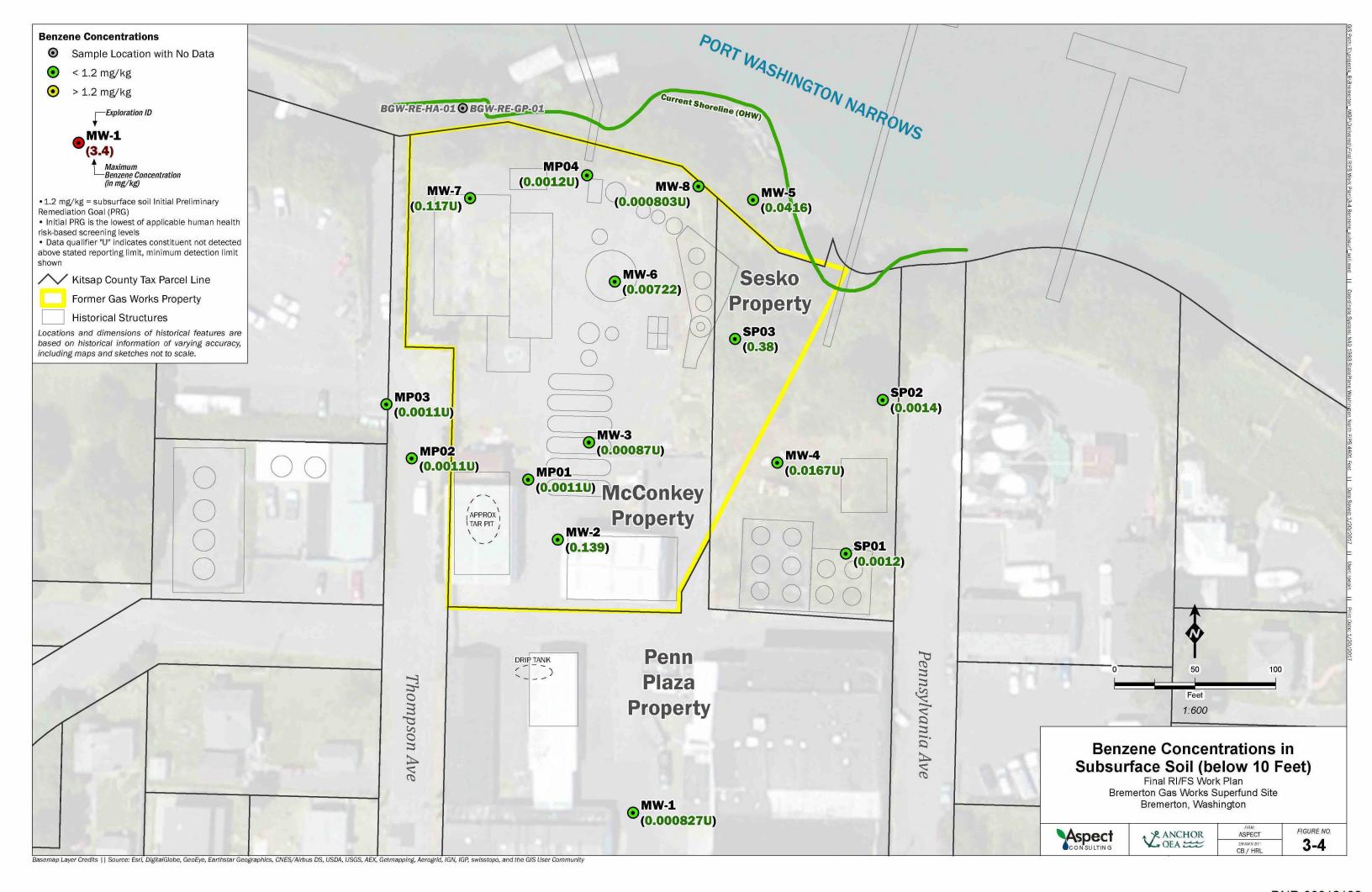


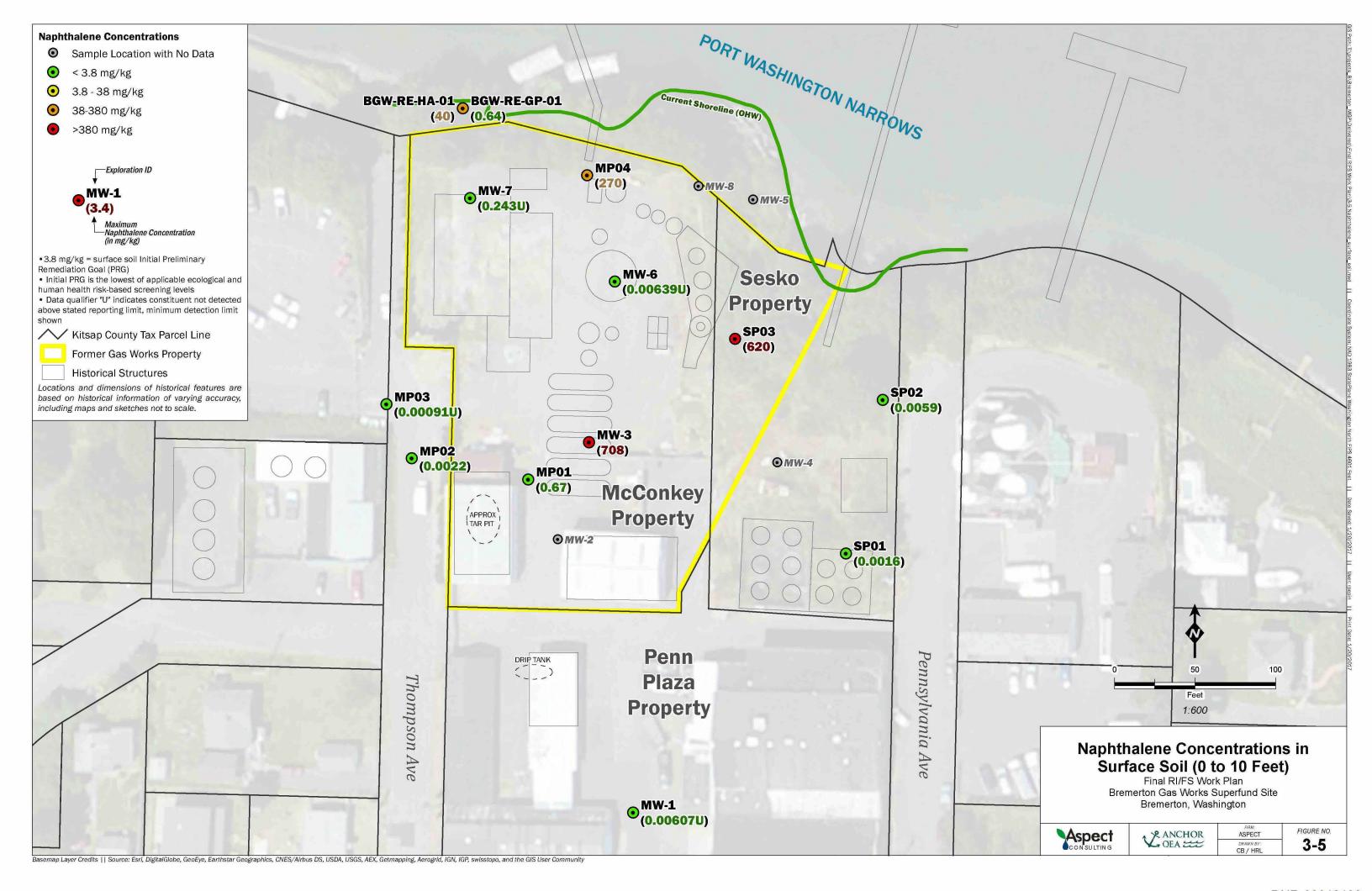


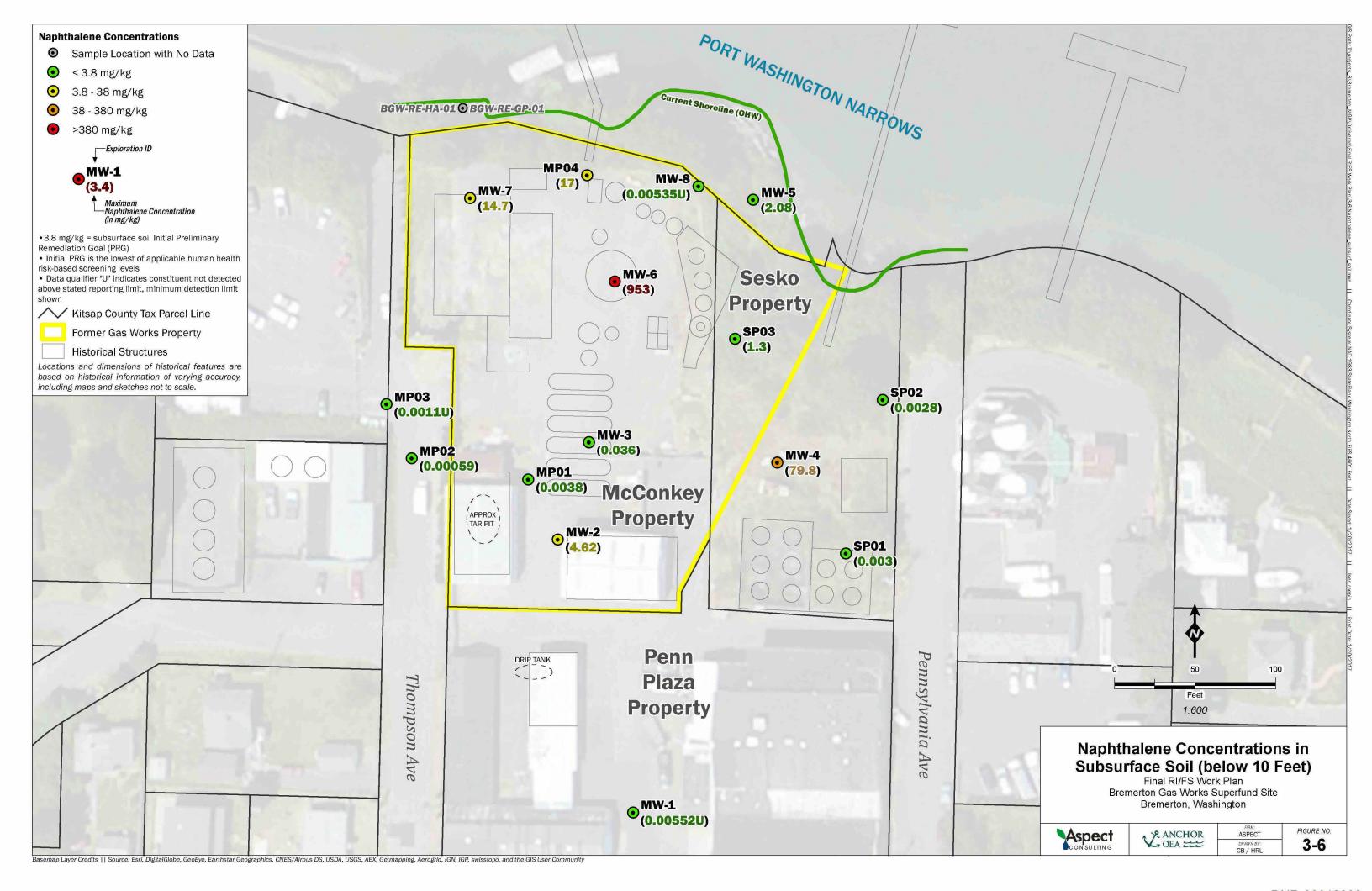


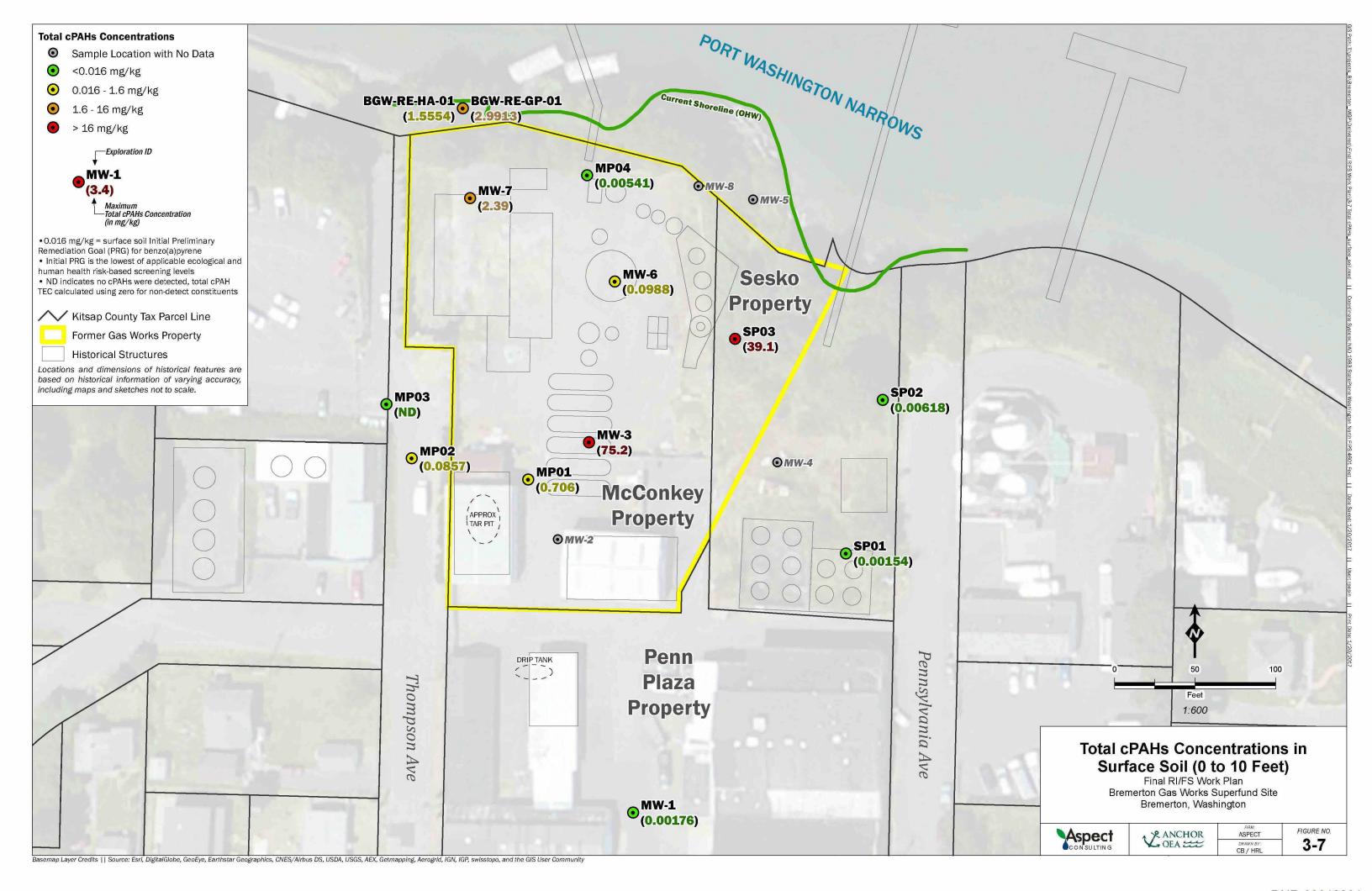


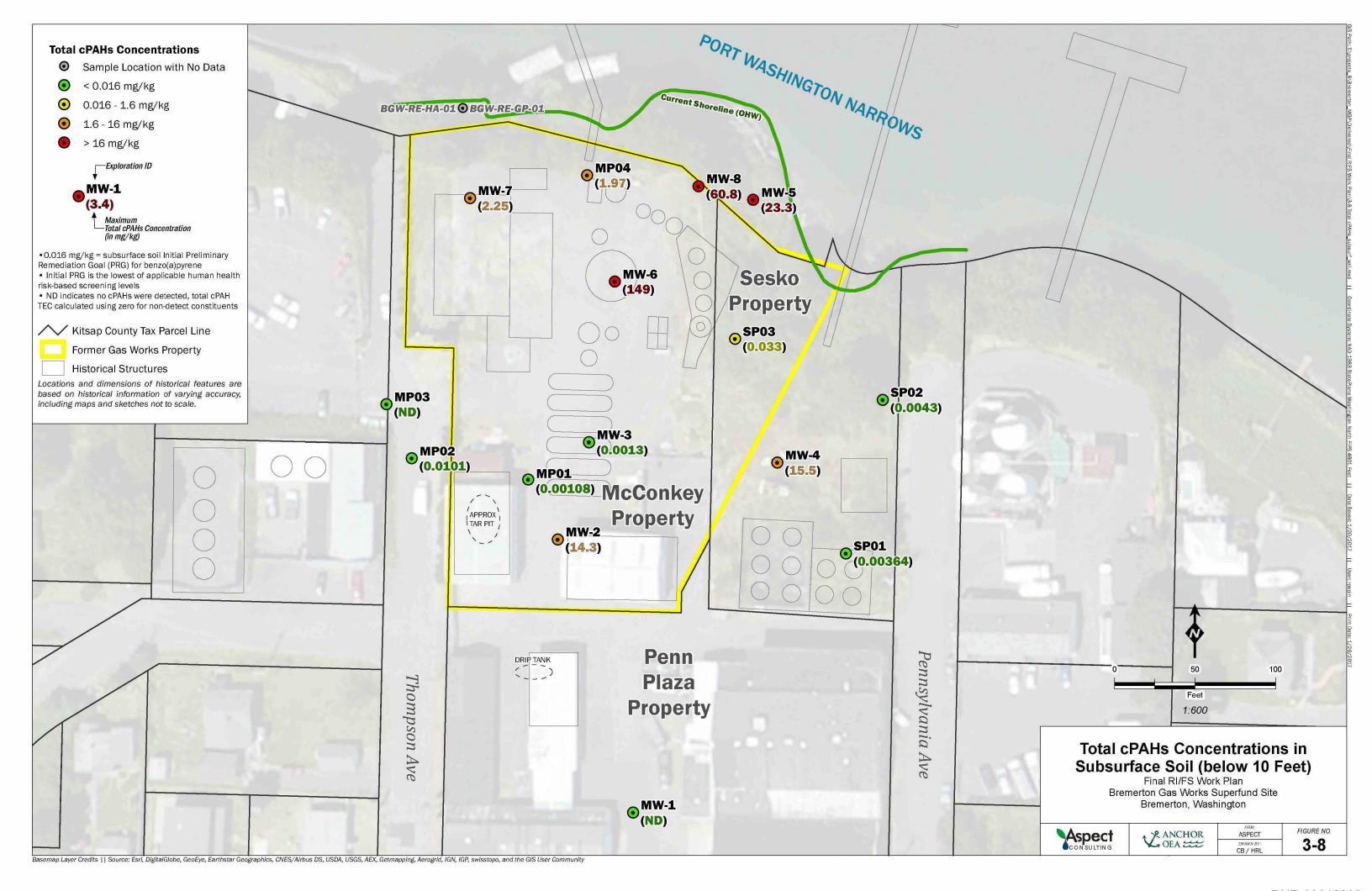


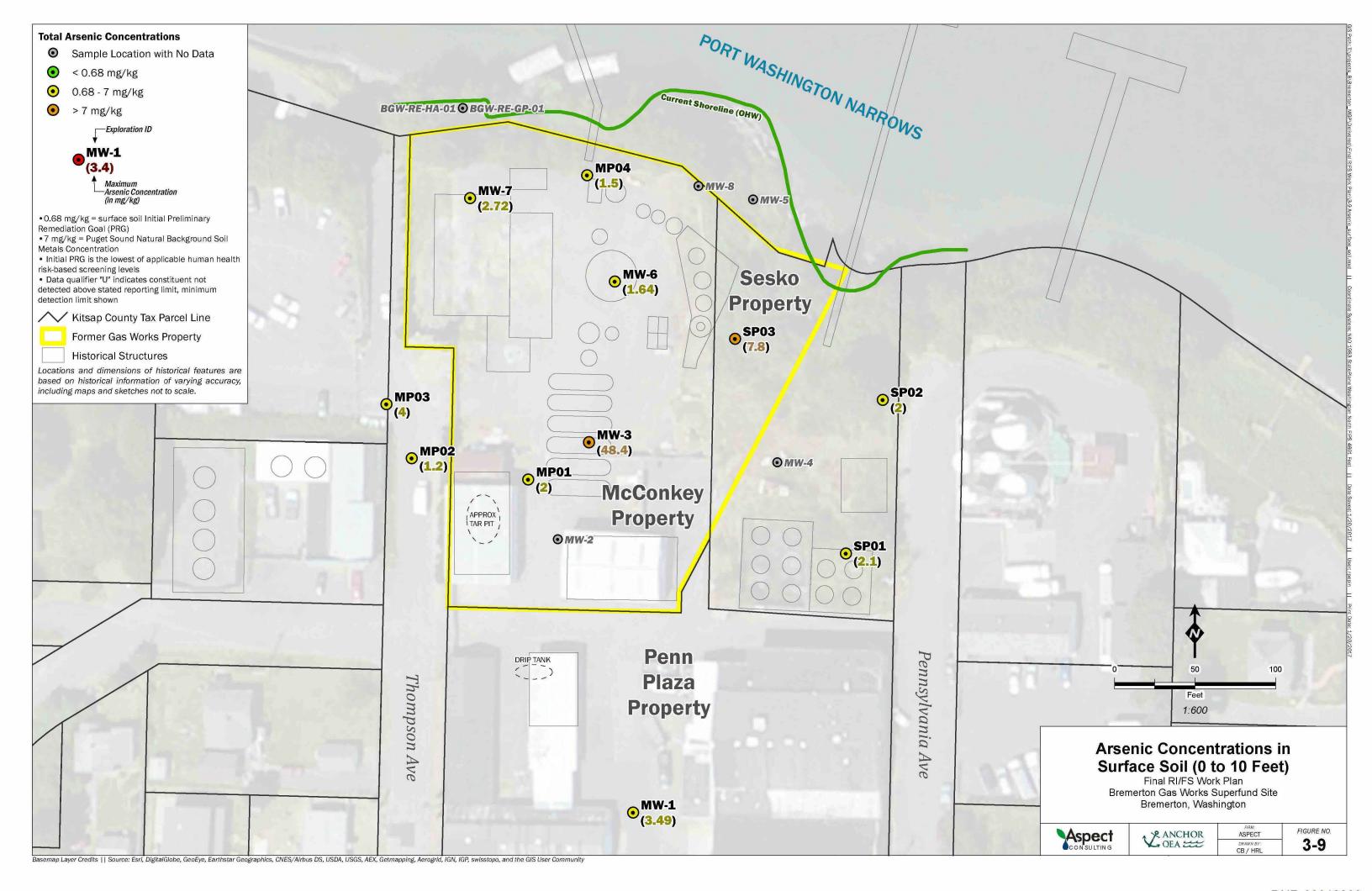


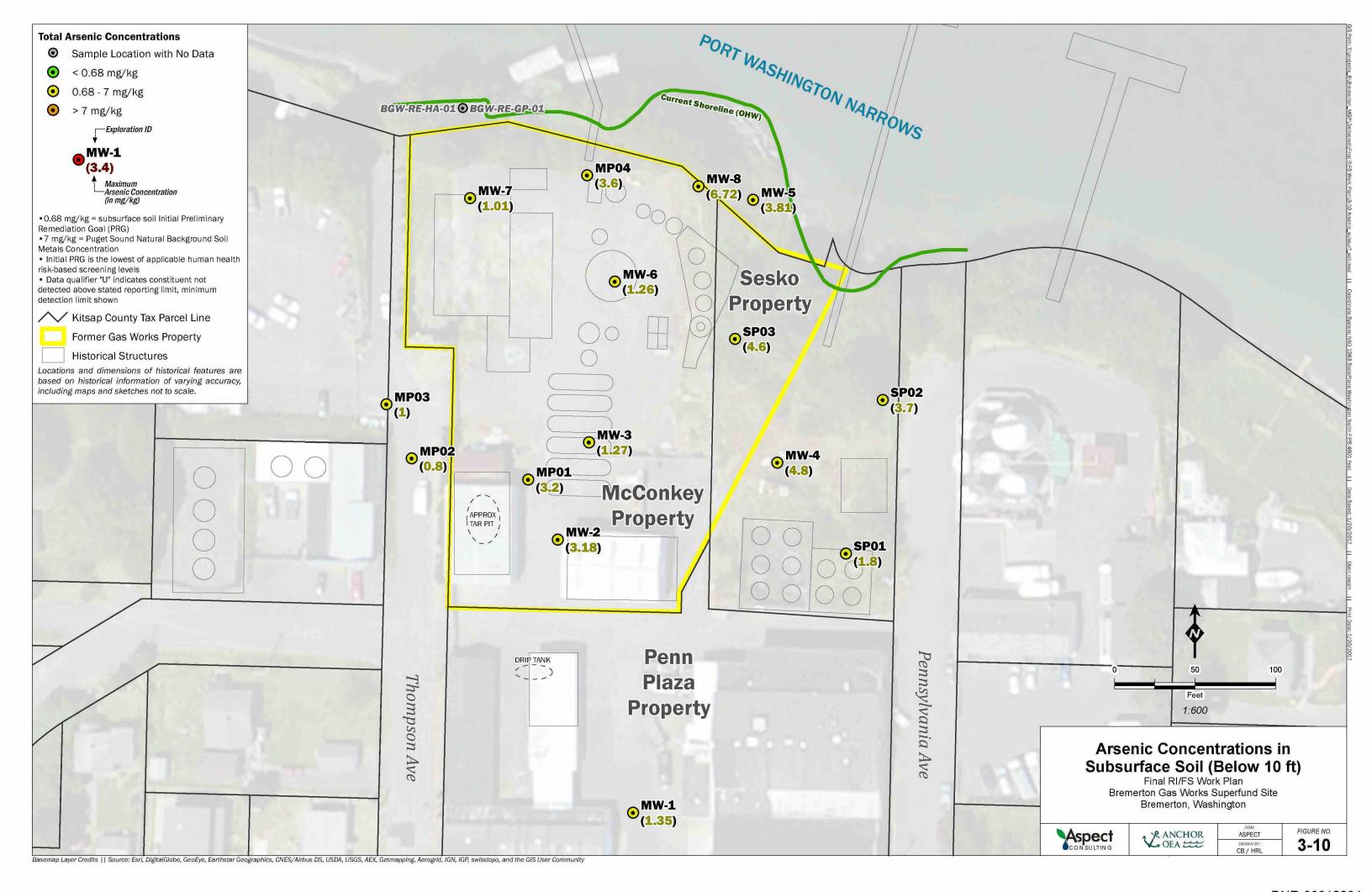


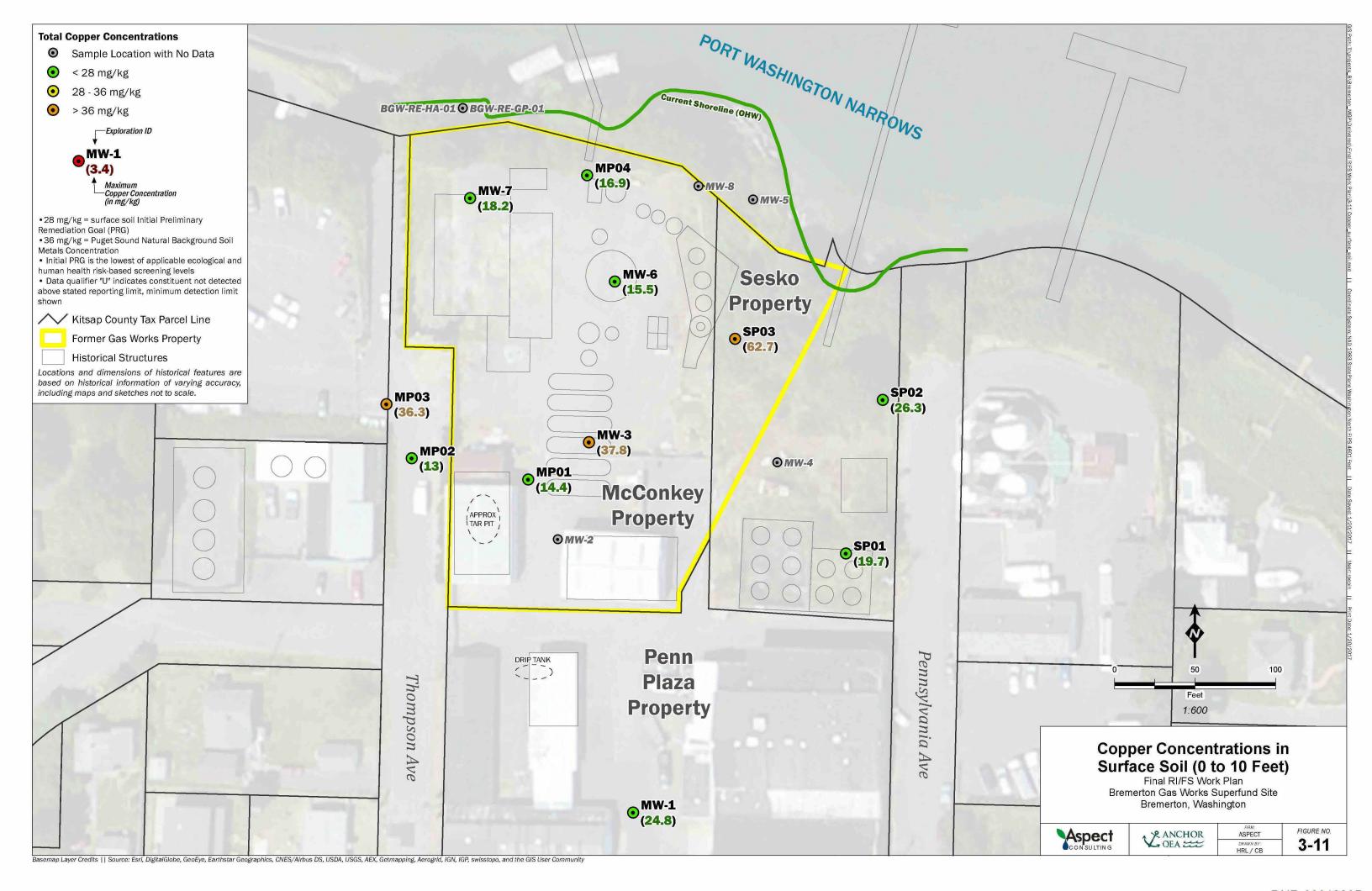


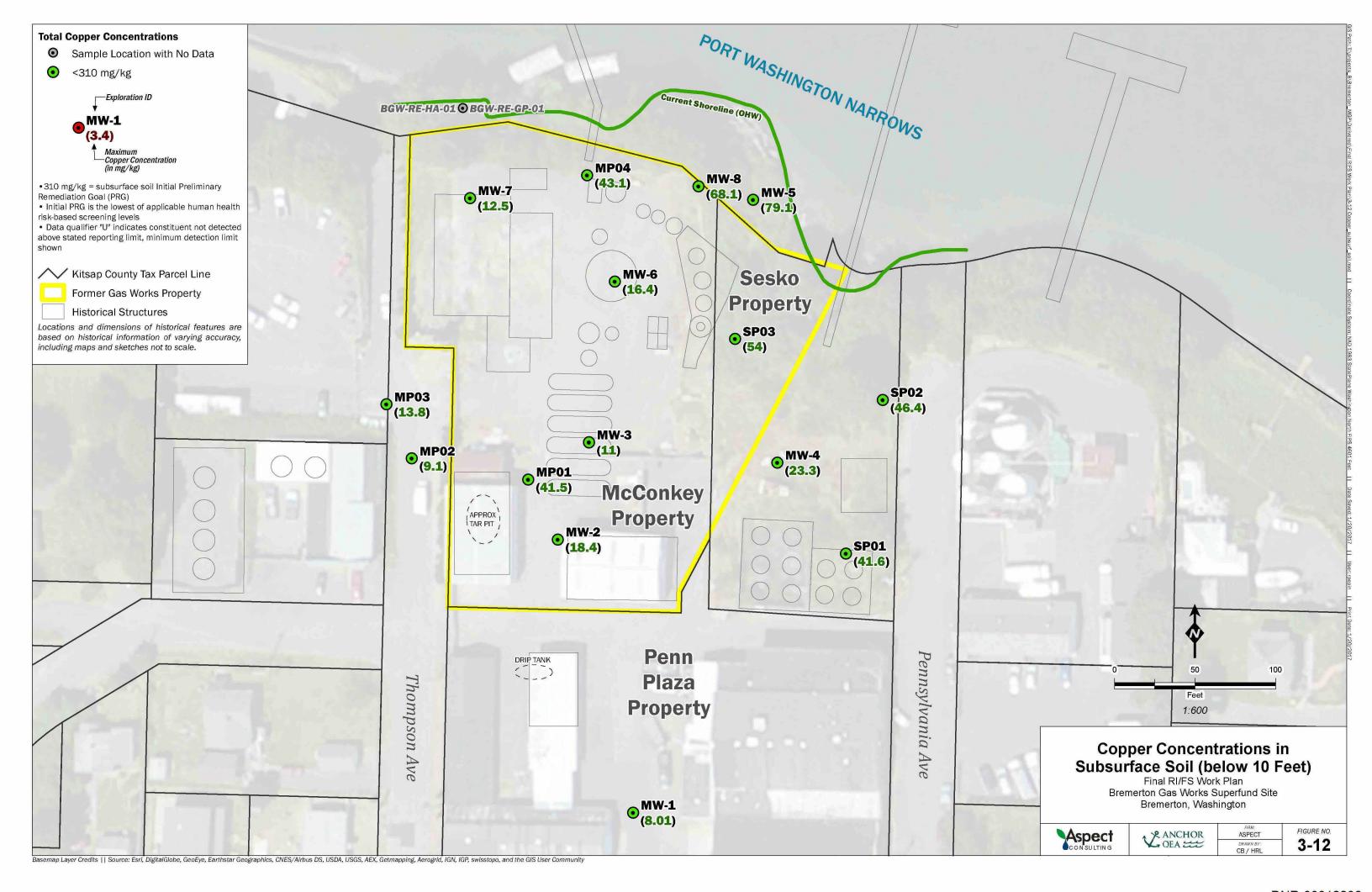


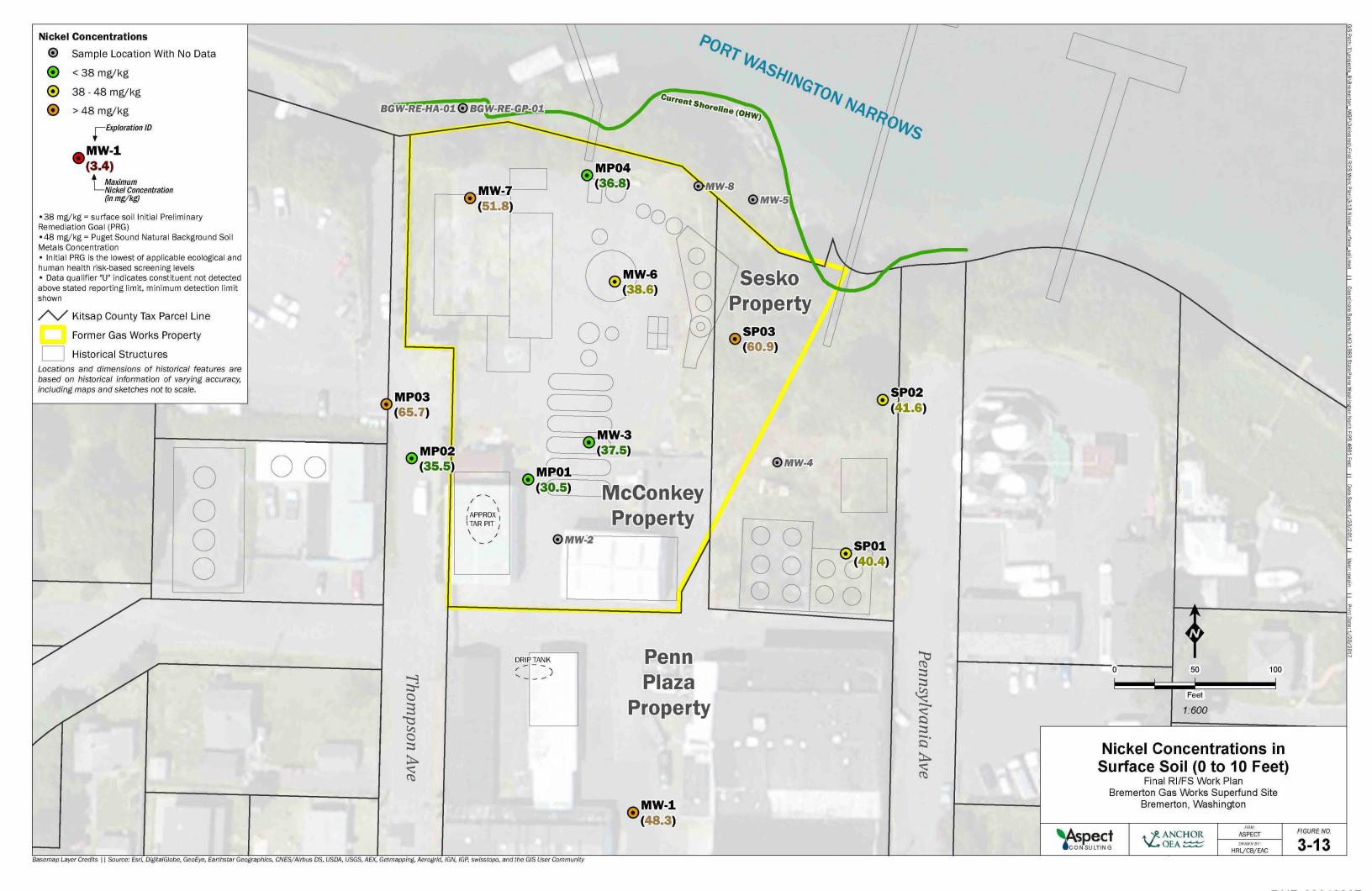


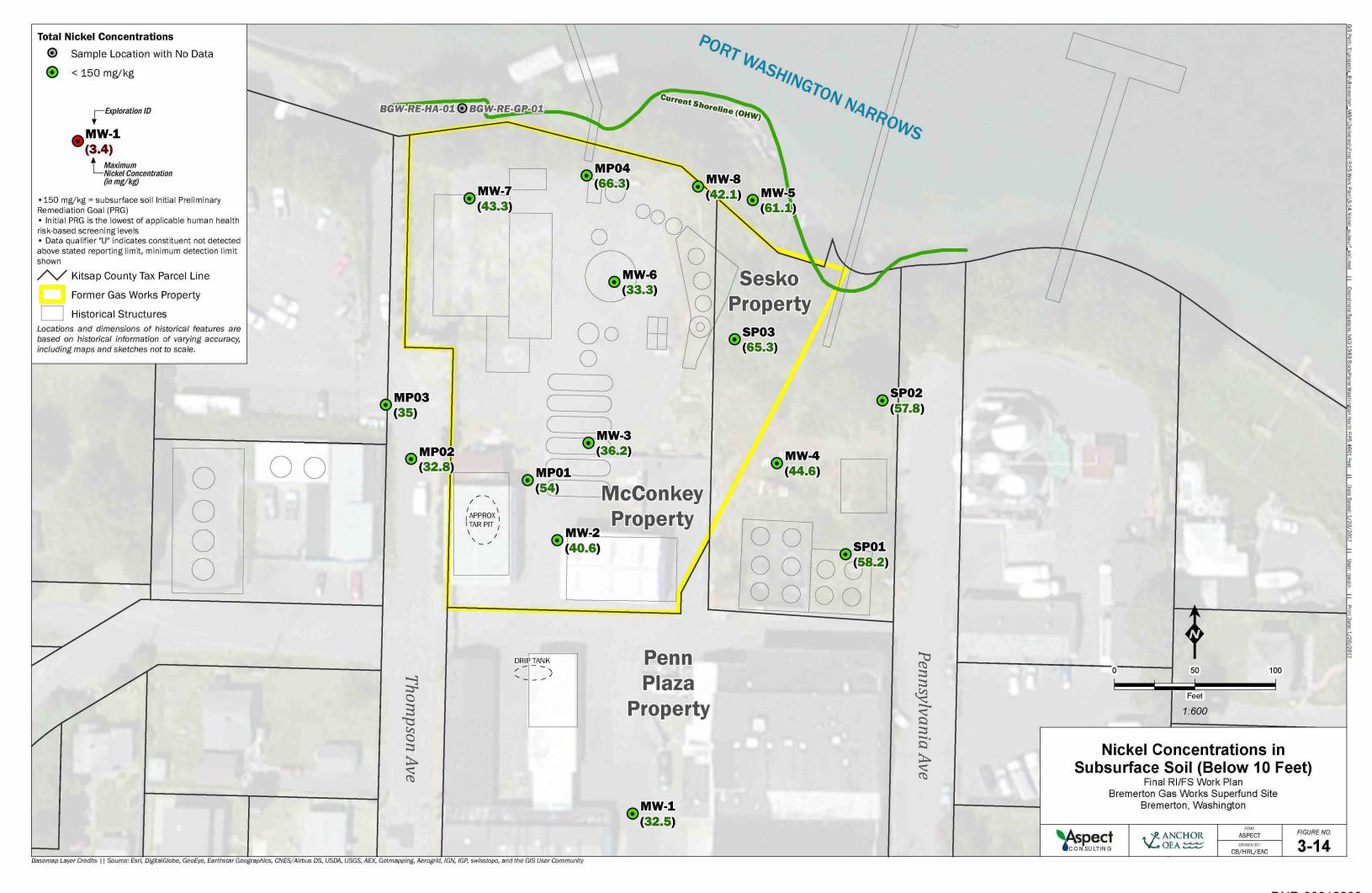


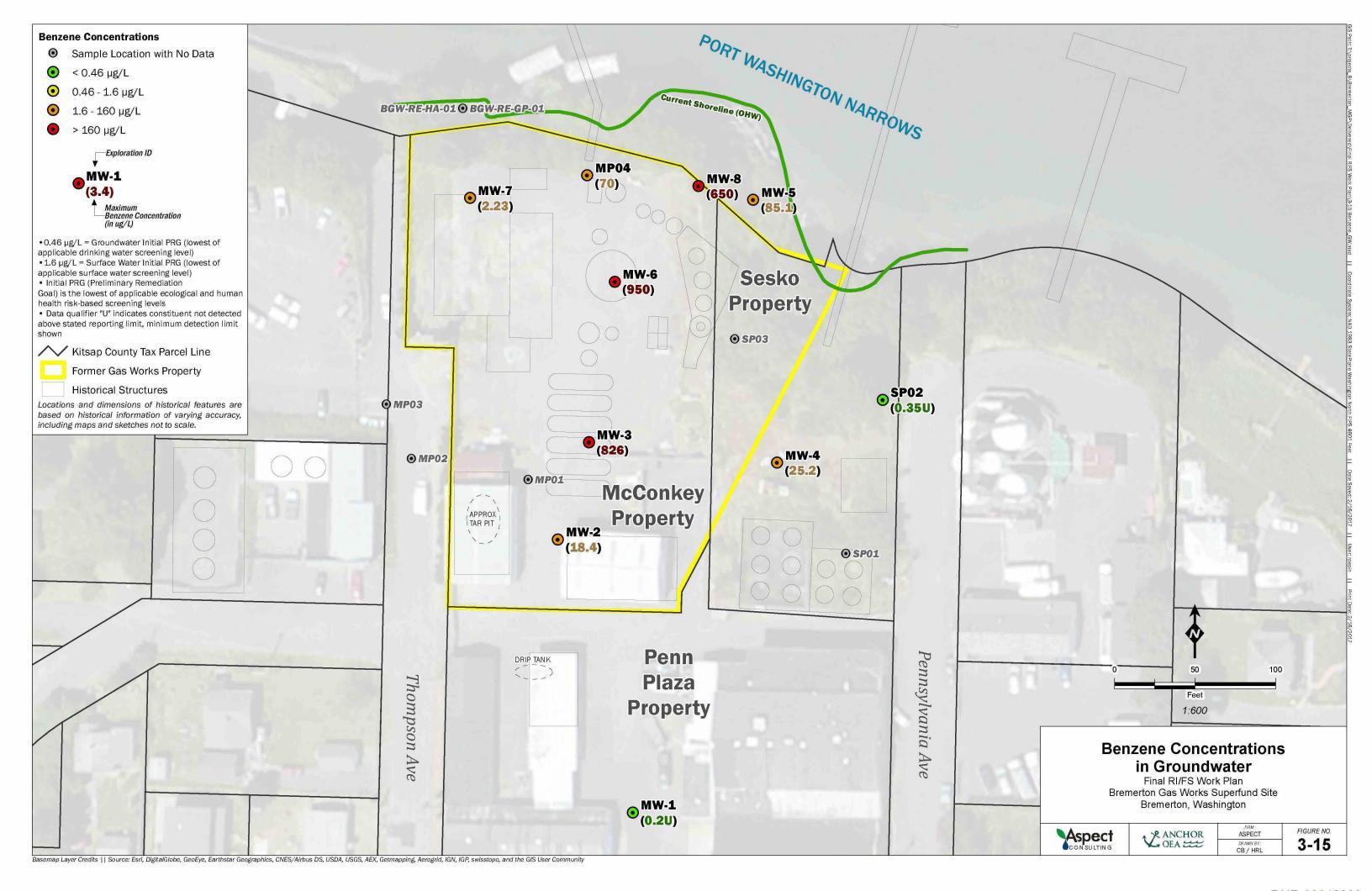


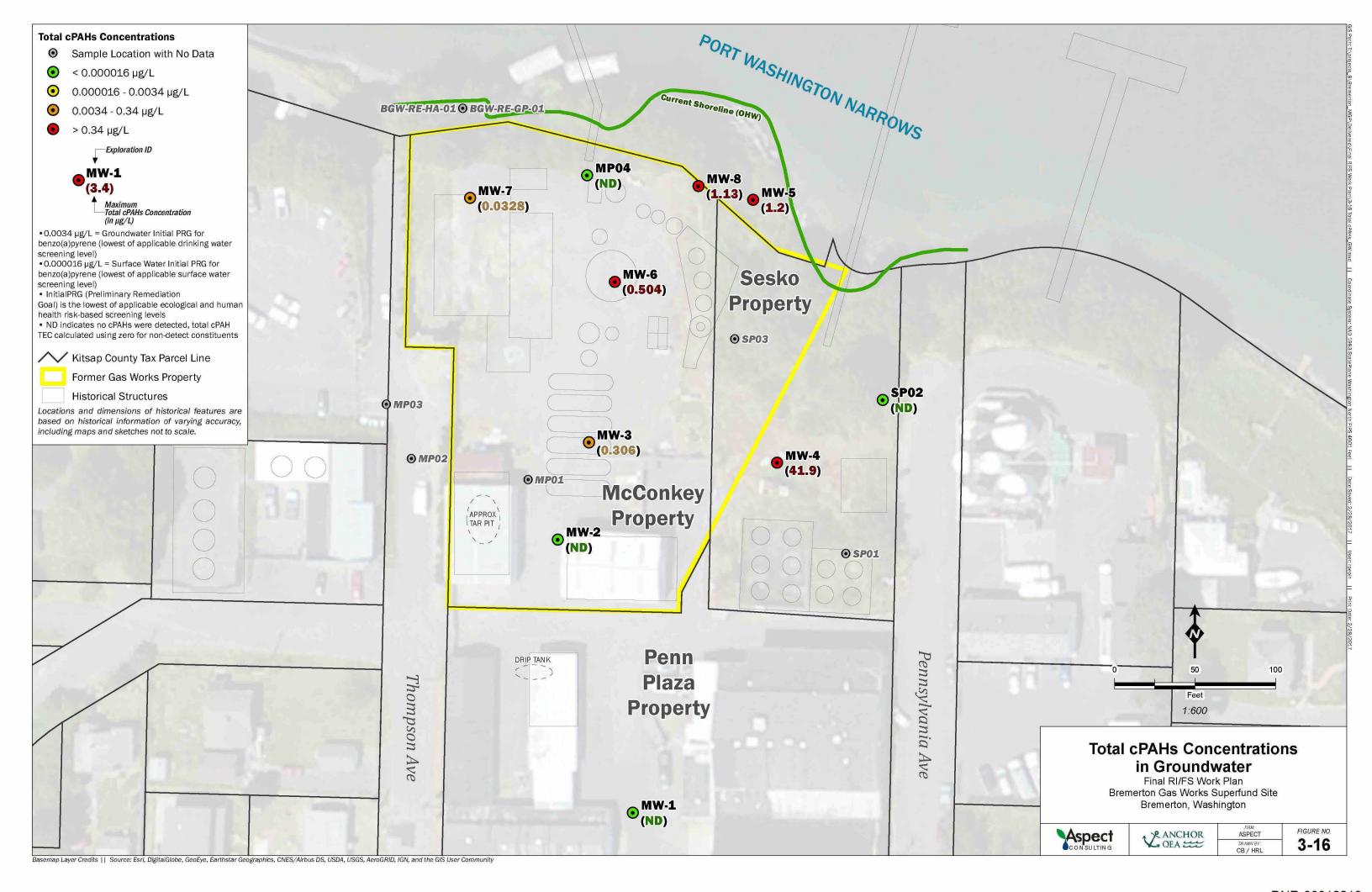


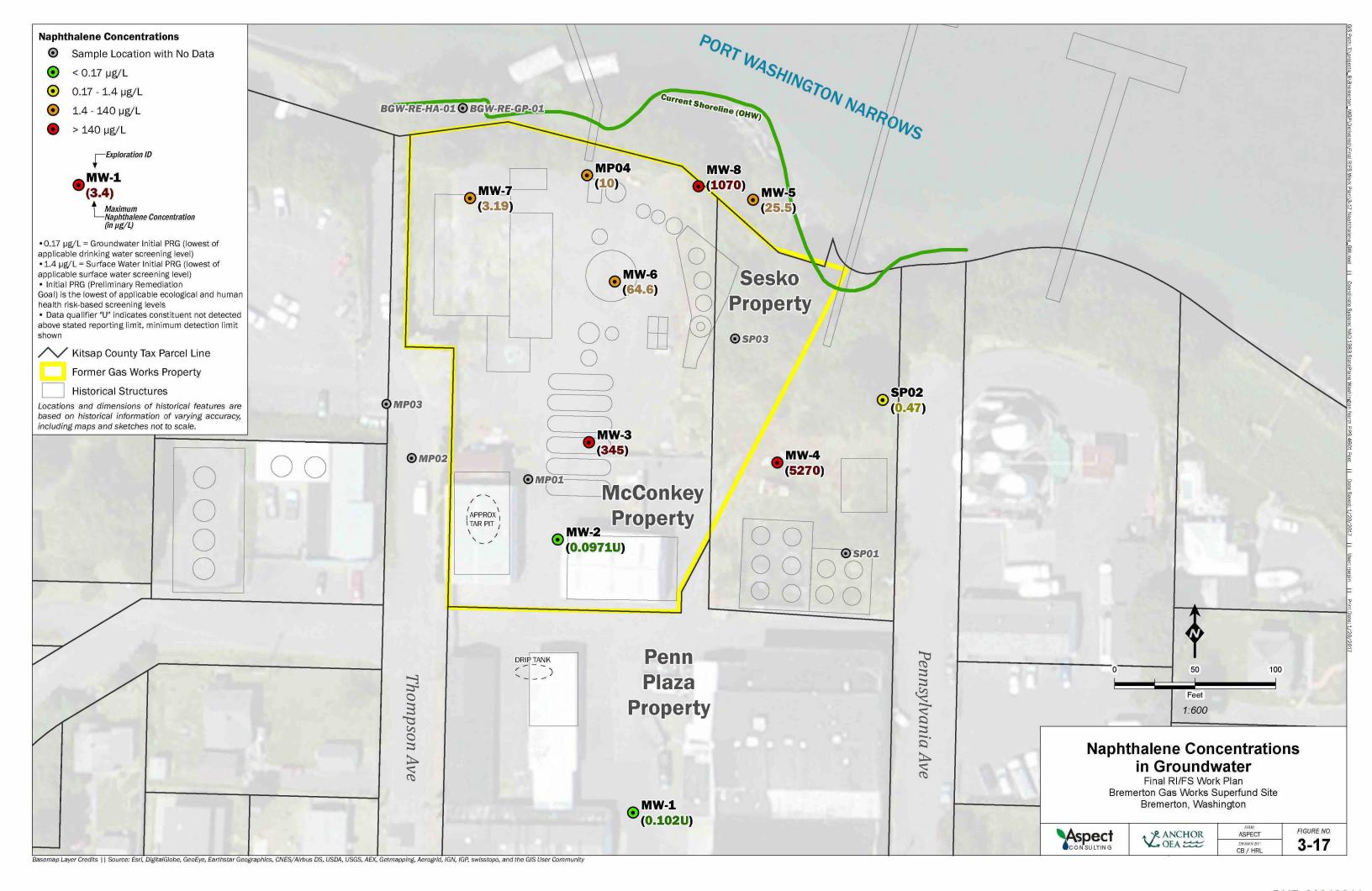


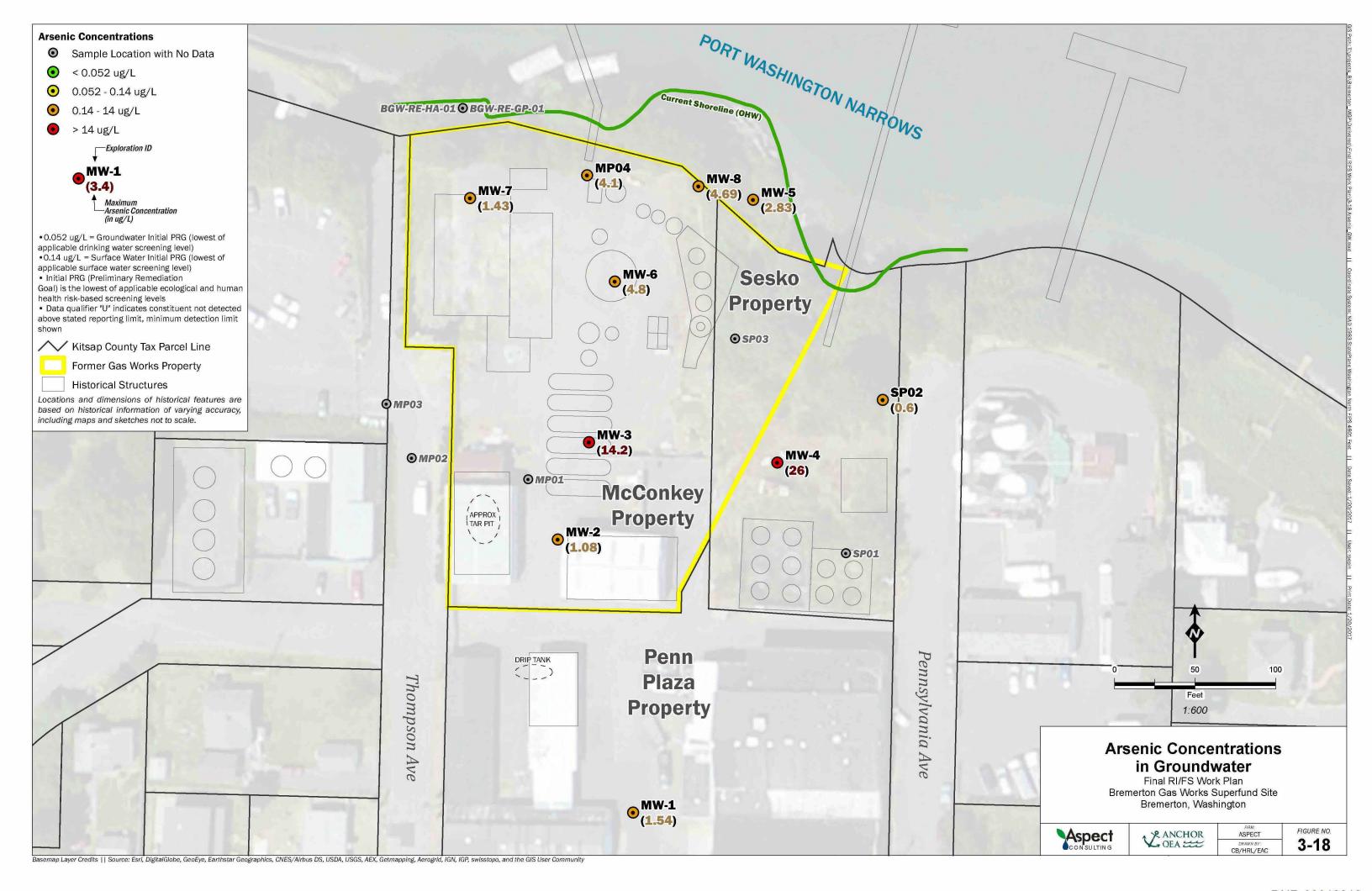


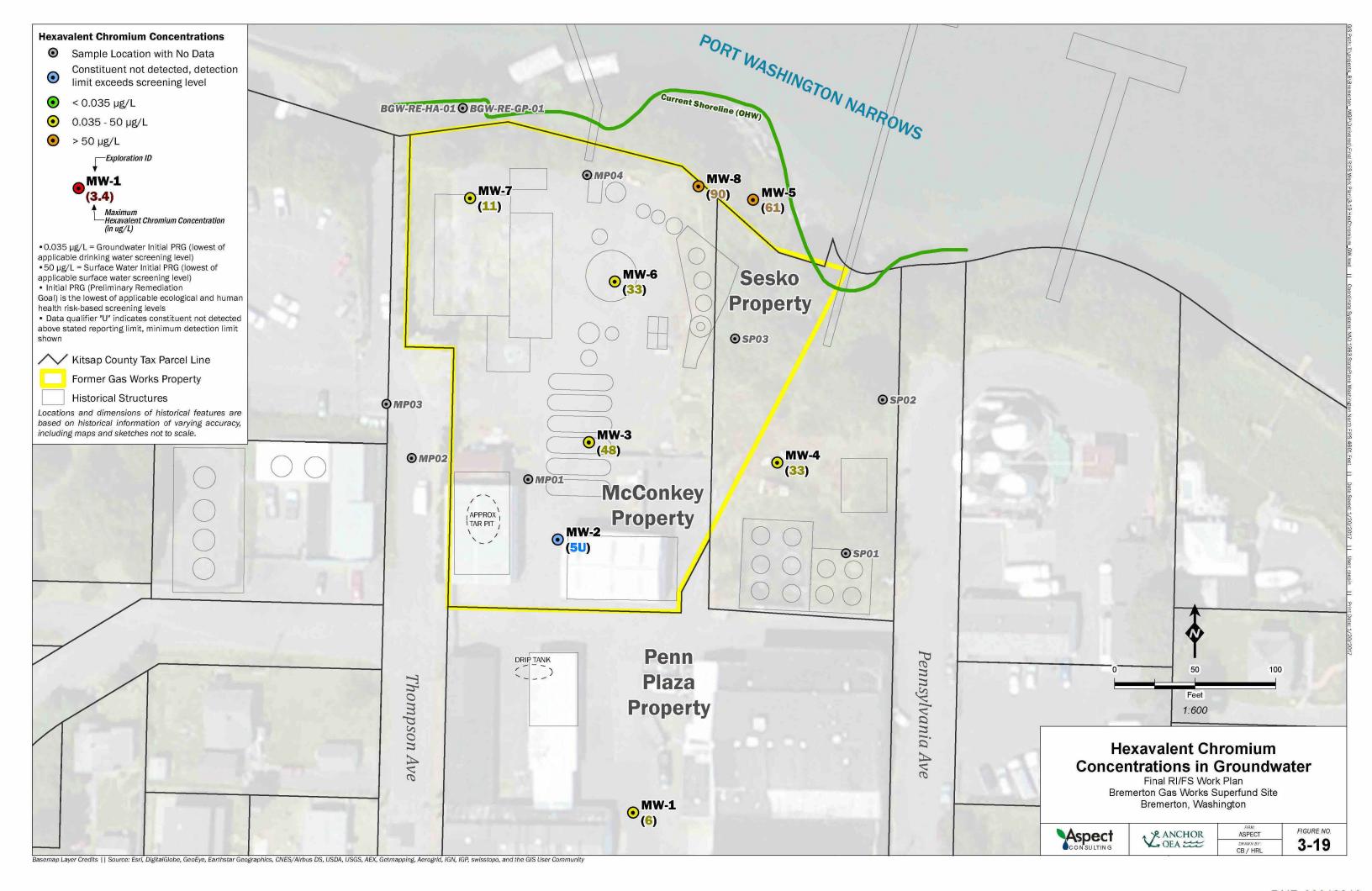


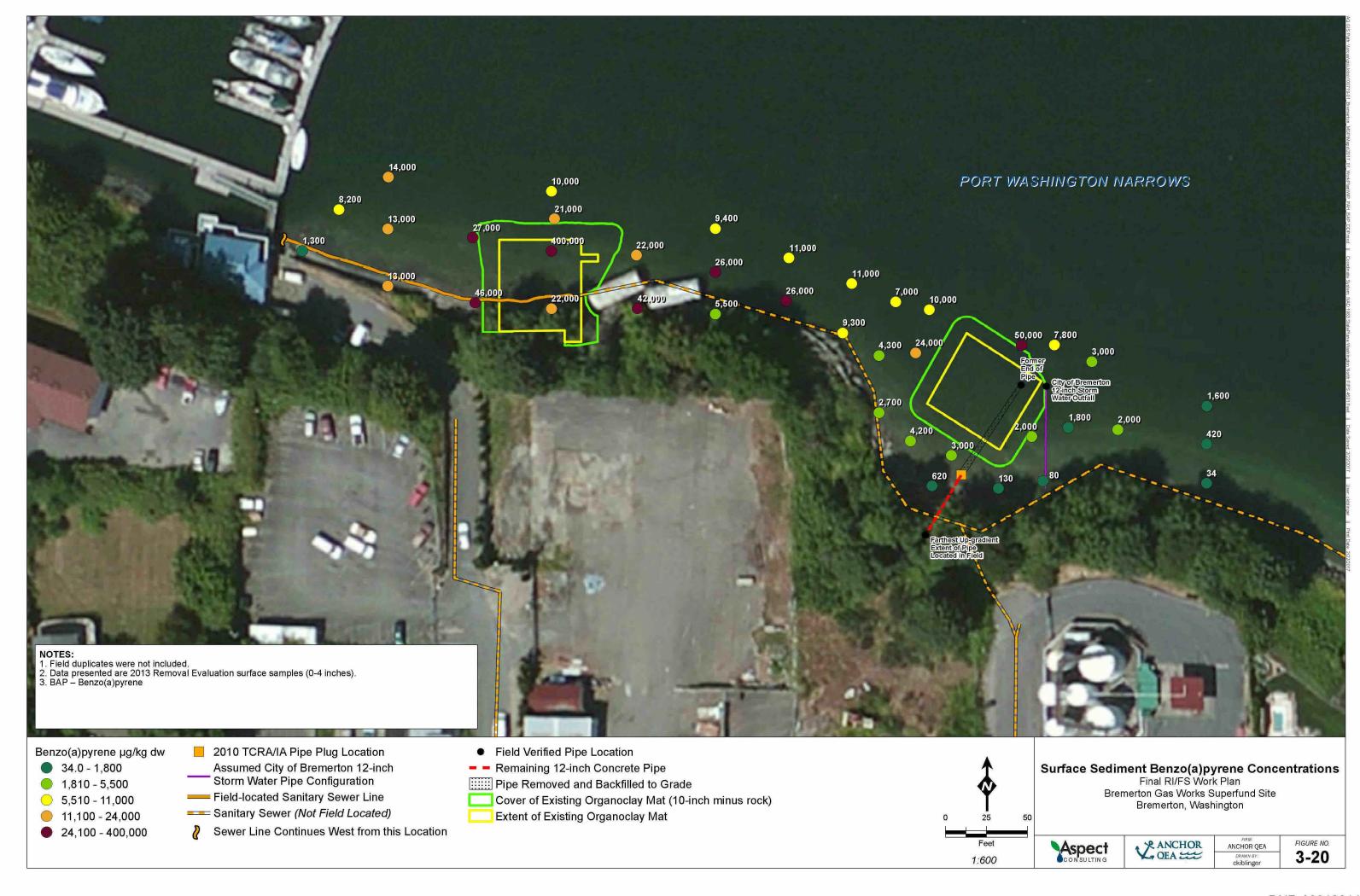


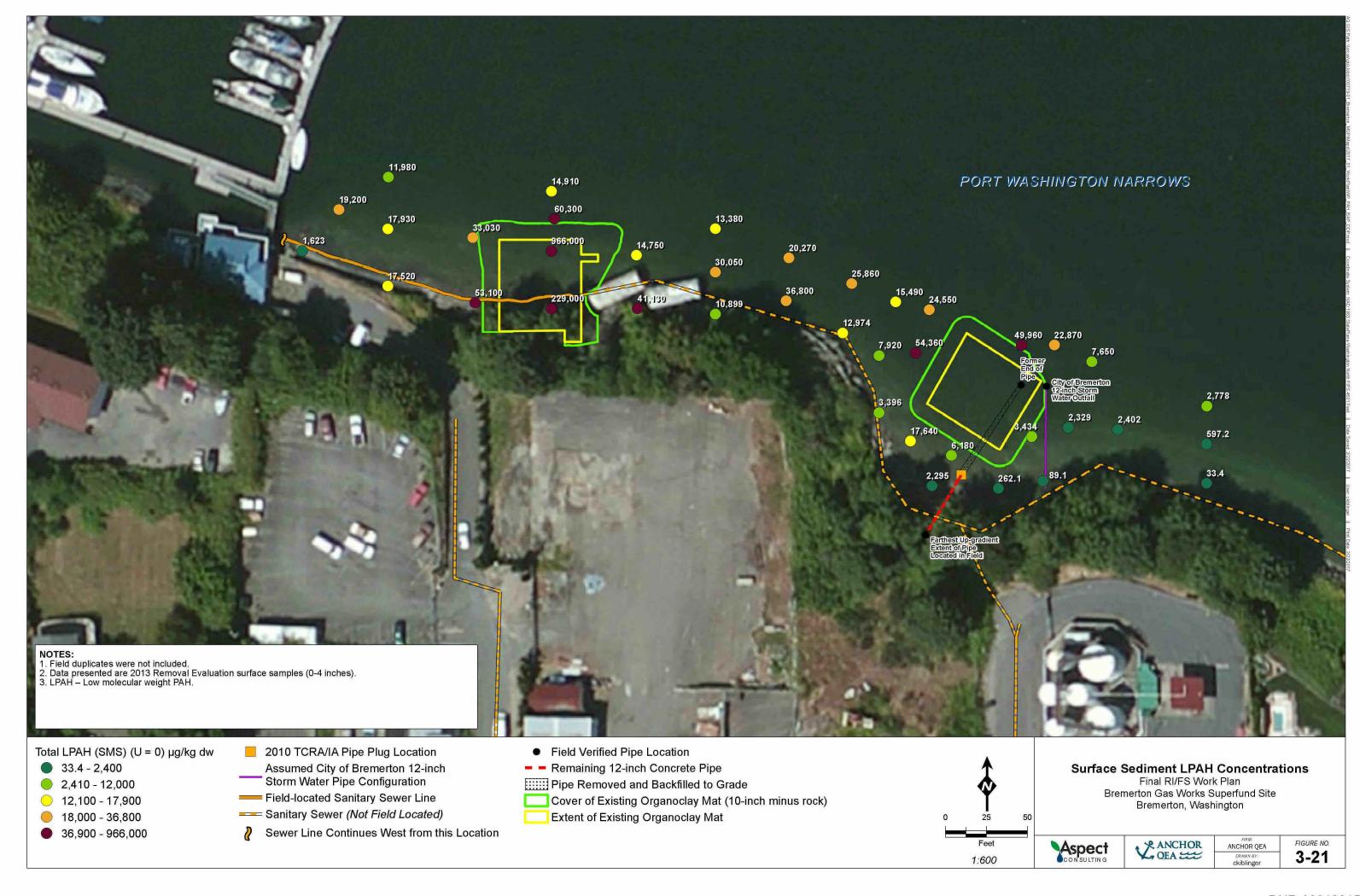


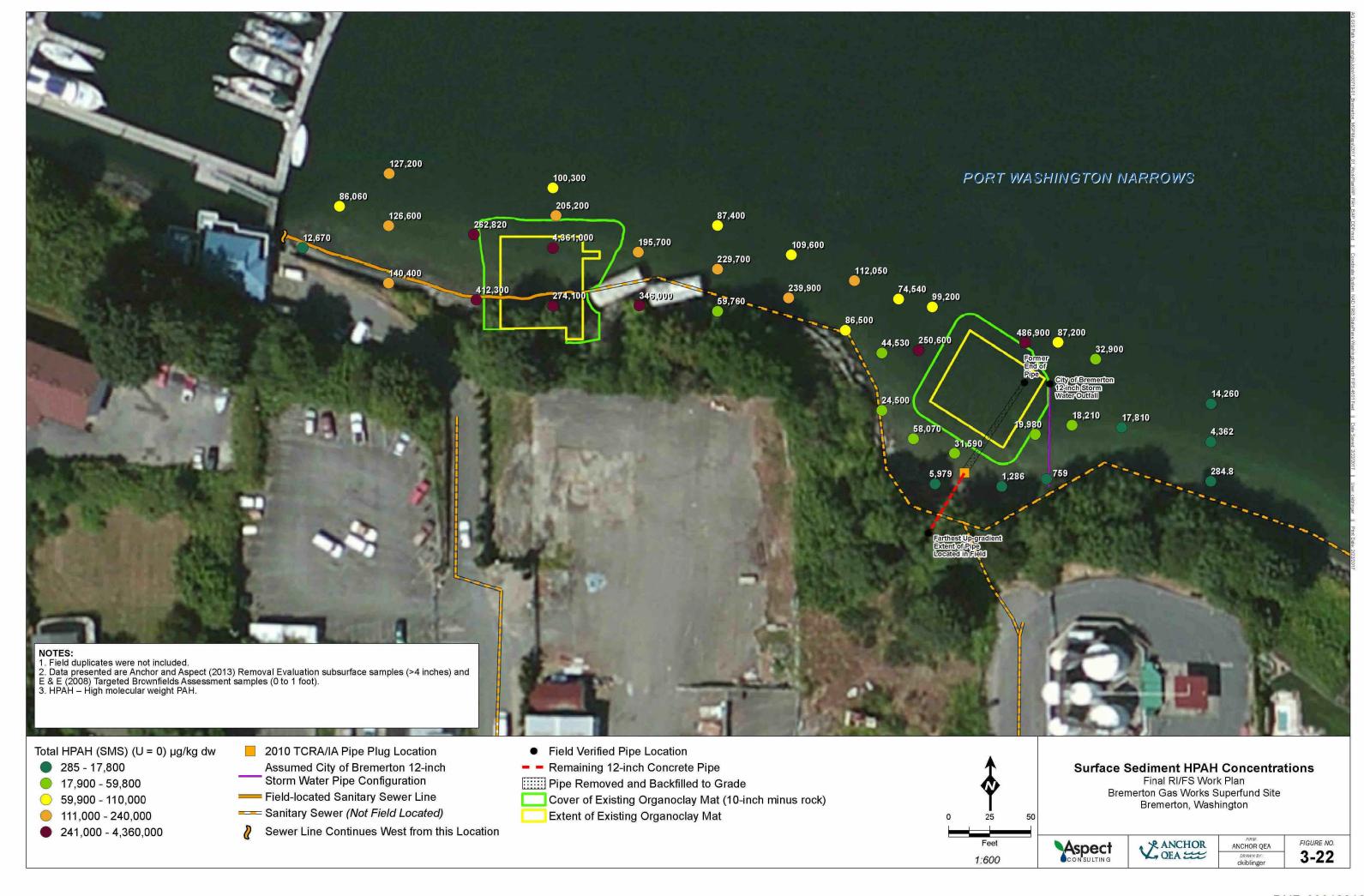


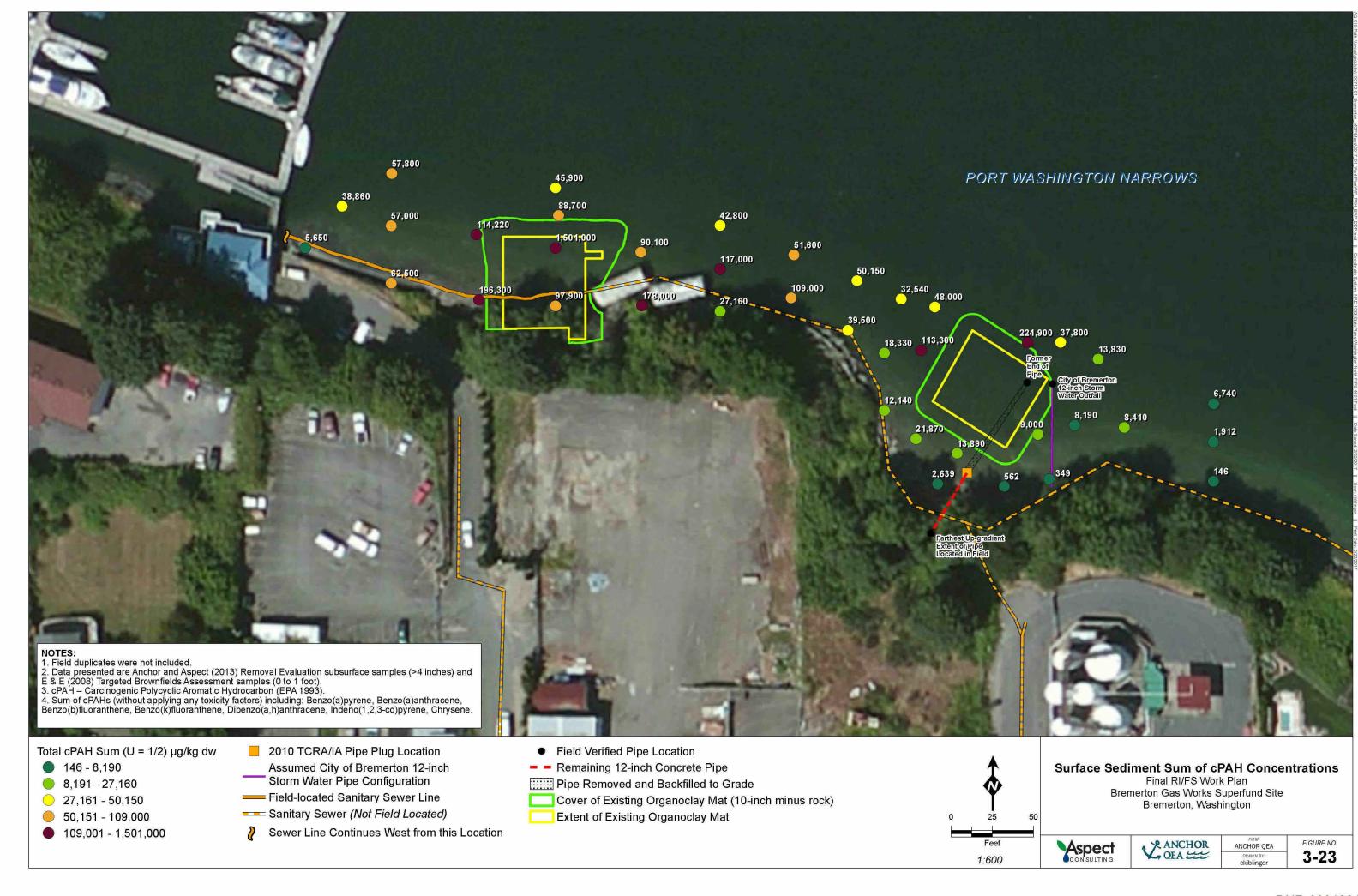


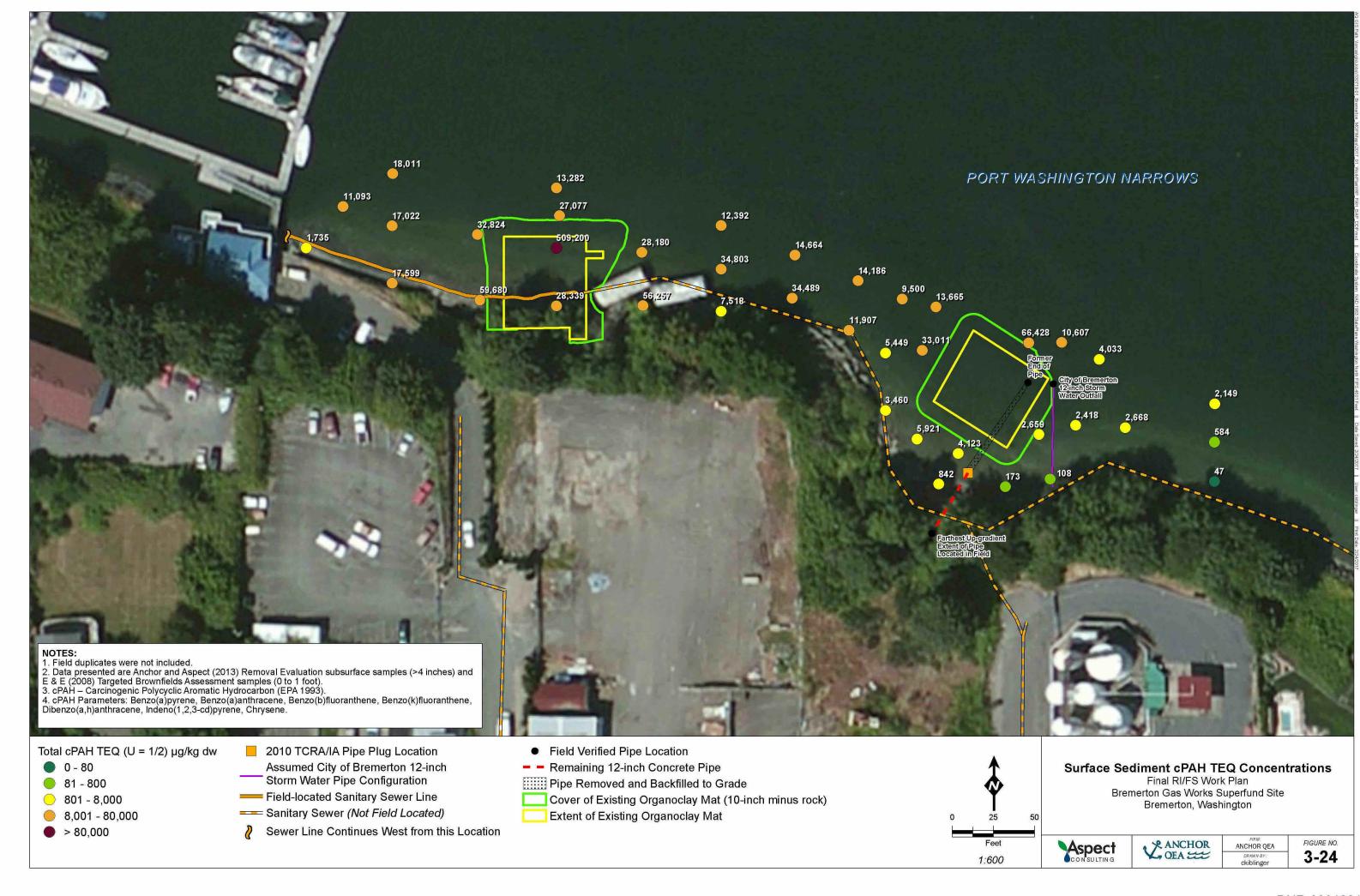


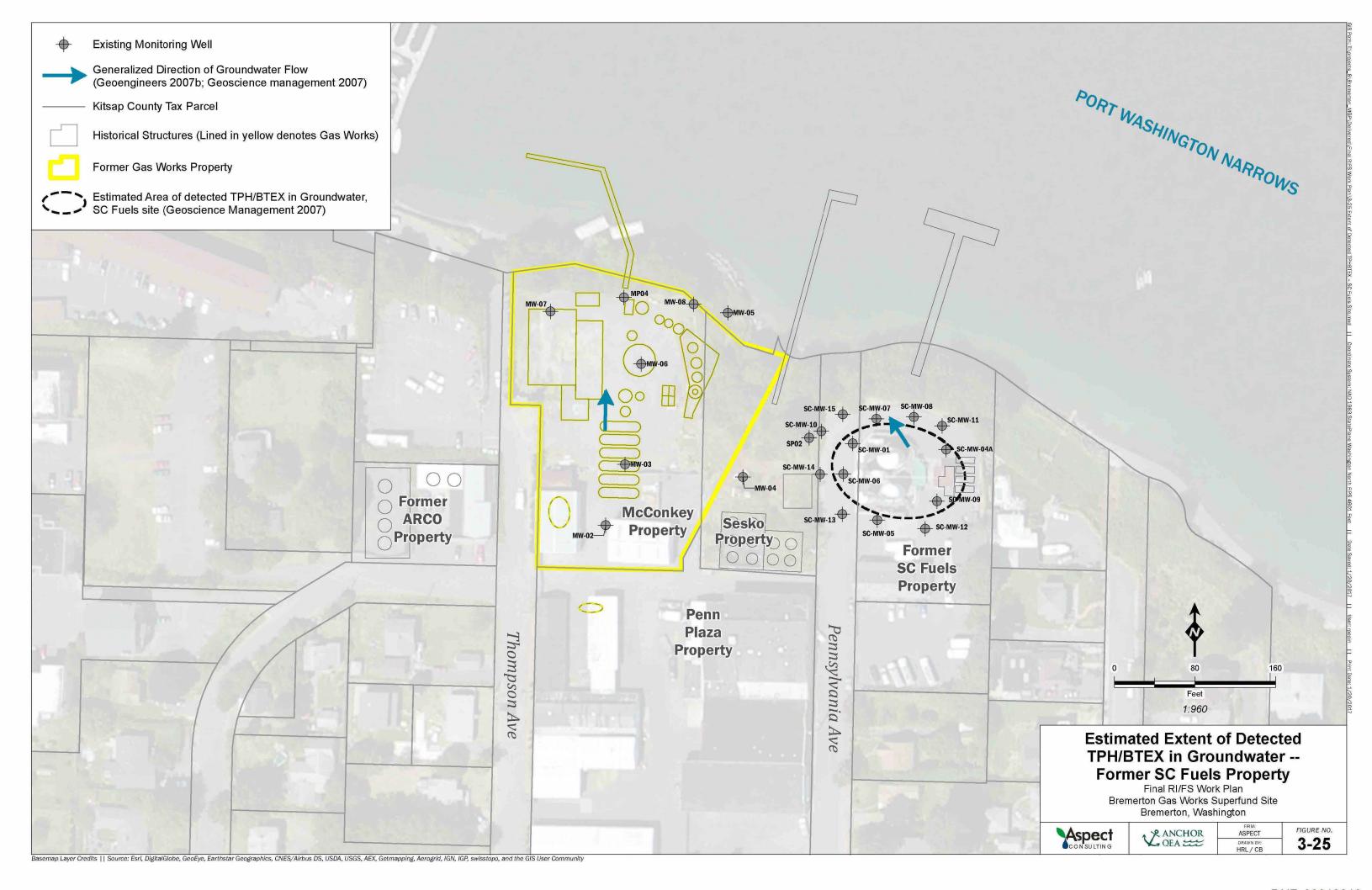


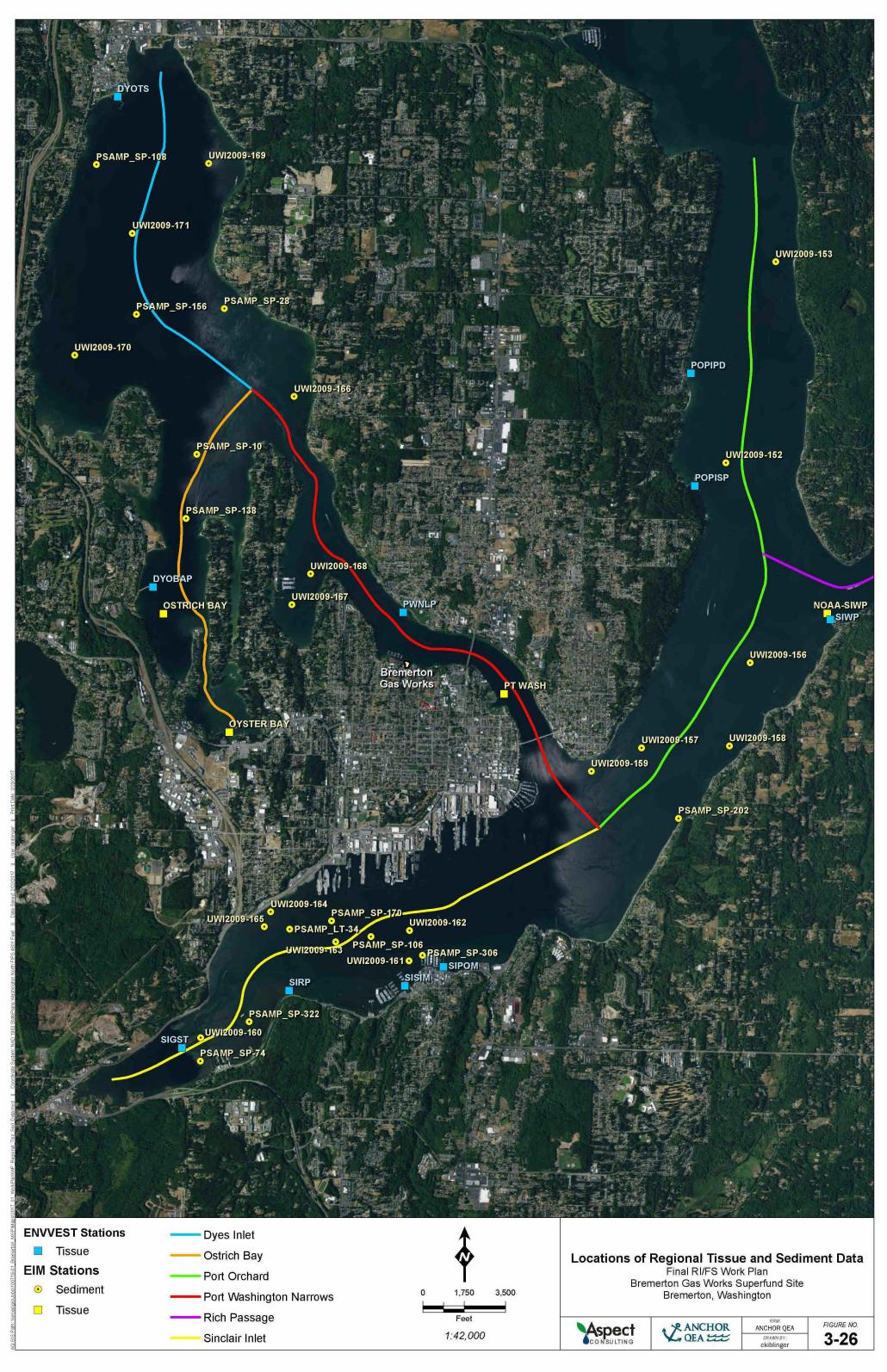


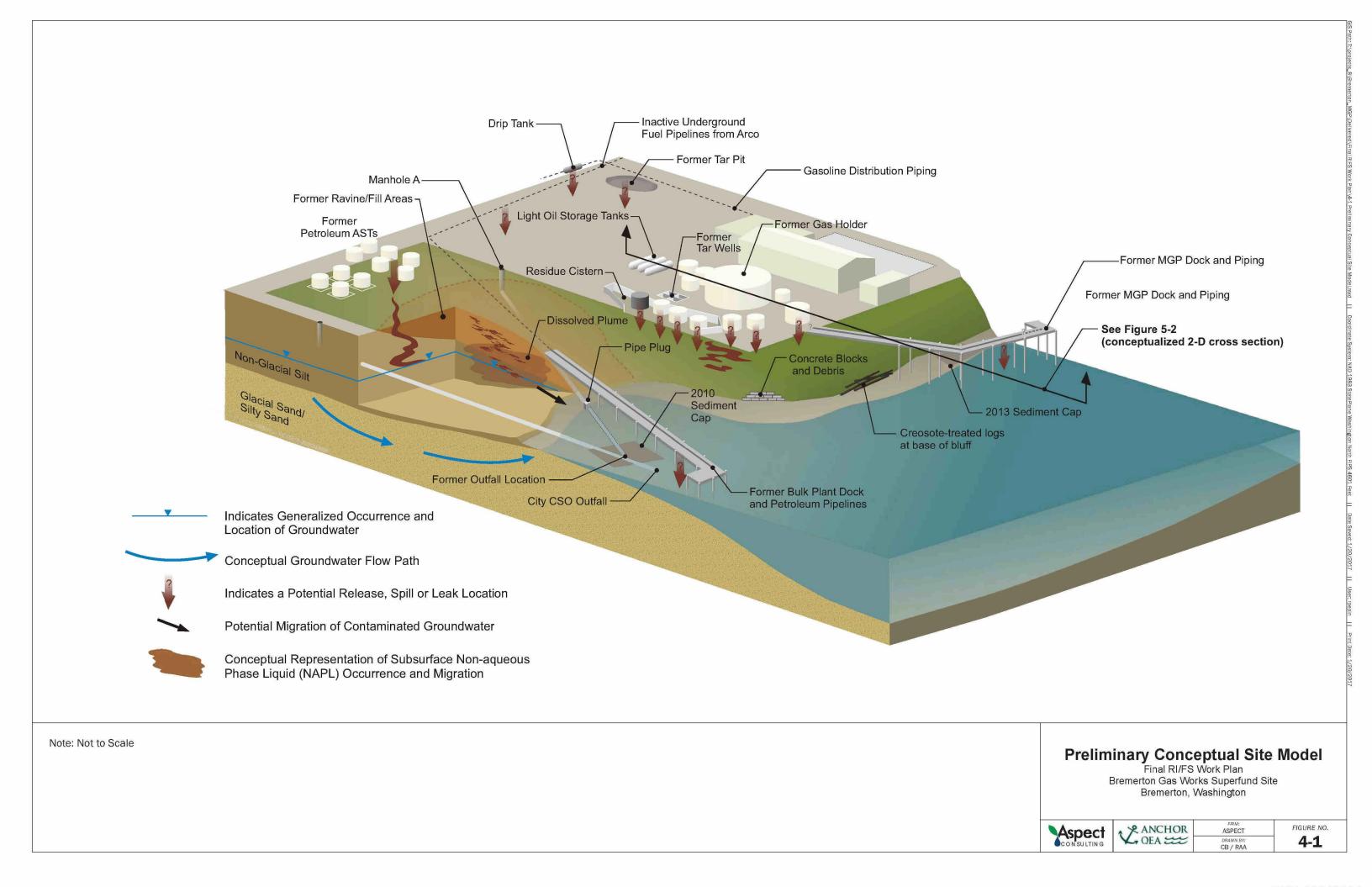


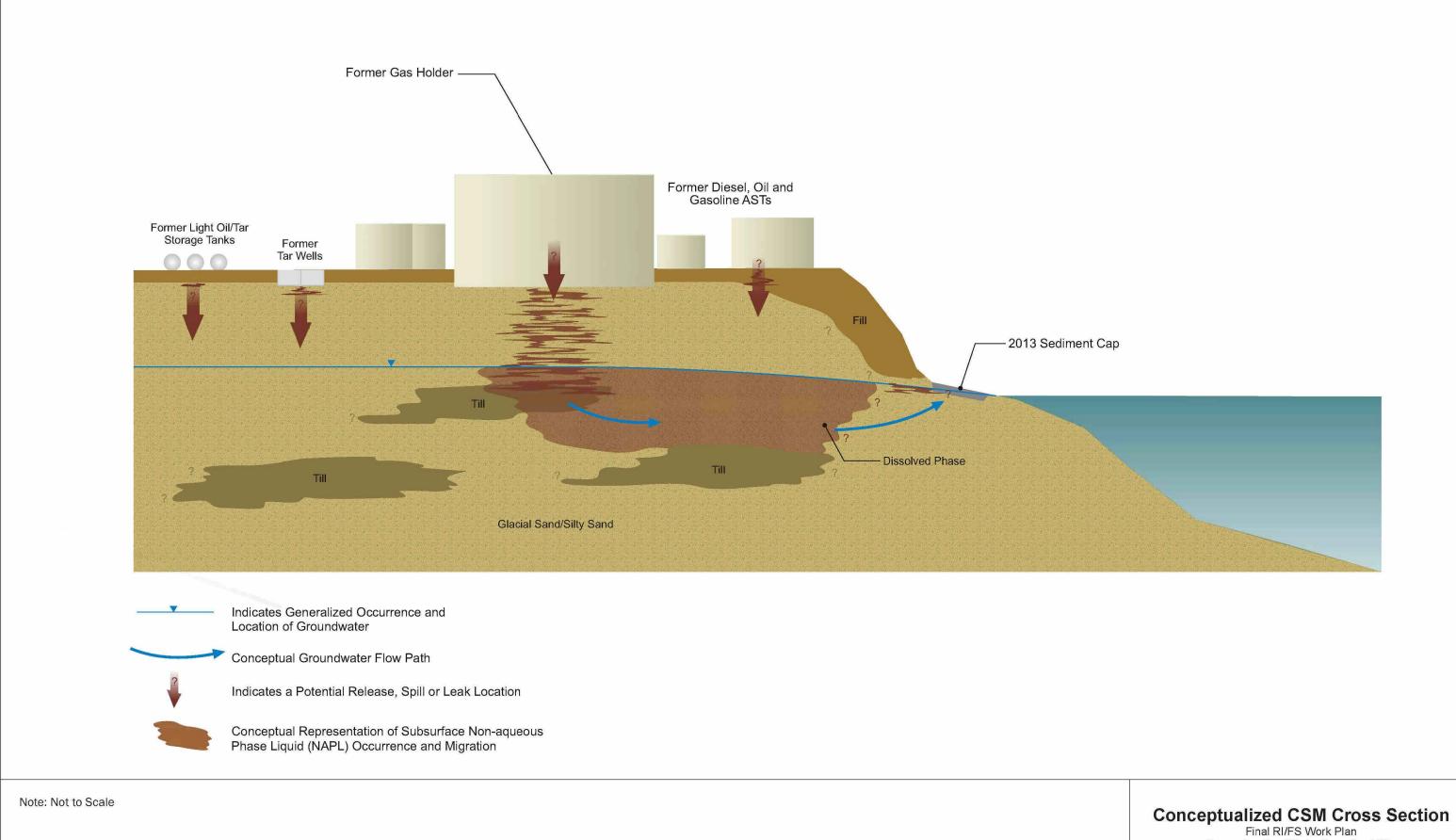










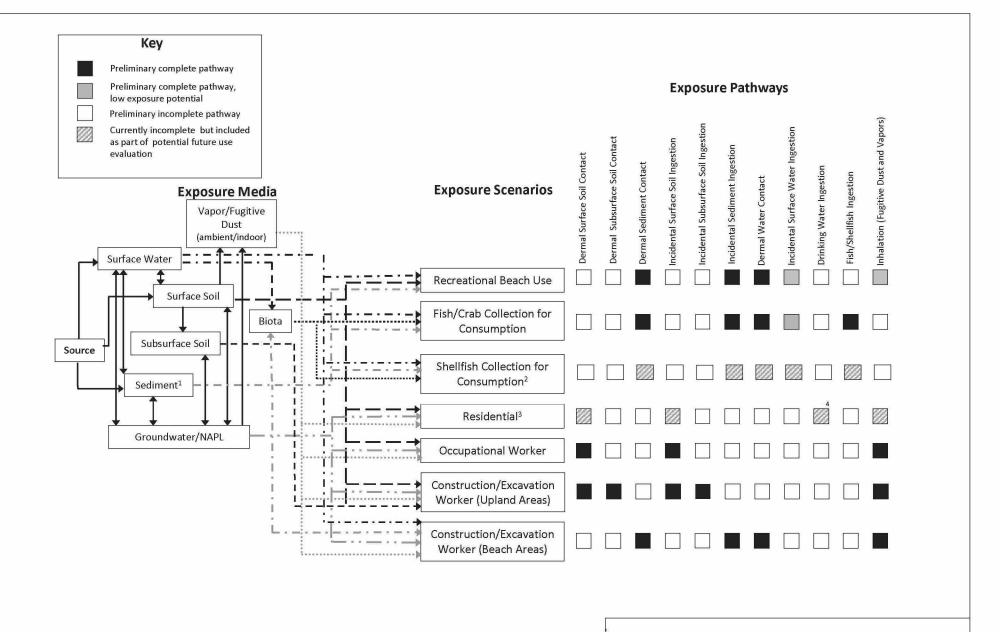


Bremerton Gas Works Superfund Site Bremerton, Washington



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FIRM: ASPECT FIGURE NO. 4-2



Notes:

- 1 Includes sediment porewater
- The portions of Port Washington Narrows adjacent to the Gas Works are currently listed as closed to shellfish harvesting (due to water quality concerns associated with combined sewer overflows and issues not related to the site) by the Washington Department of Health; however, exposures associated with shellfish harvesting will be evaluated to understand potential risks should shellfish harvest restrictions be lifted in the future.
- The Gas Works property and the adjacent properties are zoned and used for industrial uses; however, residential property exposures will be evaluated to understand potential implications should property uses be converted to residential at some point in the future.
- 4 No water supply wells are located on or near the former Gas Works; however, groundwater ingestion is retained for screening pending further evaluation of groundwater beneficial uses.

Human Health Conceptual Site Model

Final RI/FS Work Plan Bremerton Gas Works Superfund Site Bremerton, Washington





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FIGURE NO. **4-3**

